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MASTER THESIS DRAFT

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1 Introduction

Ultracold atoms present an extremely powerful tool in quantum reasearch. At the scale of nanokelvins thermal effects no longer destroy the otherwise fragile quantum systems. Thus, quantum phenomena, otherwise only known from theory, can be studied directly. Atomic physics experiments with such ultracold, quantum degenerate gases offer some unique features: (i) a wide range of Hamiltonians can be mapped to the systems making experiments highly customizable. This can in part be attributed to (ii) the very high degree of control achievable through the manipulation of external fields. Through this, the cold atoms can be trapped and manipulated using magnetic and optical traps. Furthermore, the collisional properties of the atoms can be tuned through magnetic Feshbach resonances, and the properties of the gas can be probed through interactions between internal energy levels of the atoms and laser light [16, 2].

In the regime of such cold temperatures many-body phenomena such as Bose-Einstein condensation takes place, where the ground state of a system gains a macroscopic populations. Ever since the first realisation of a Bose-Einstein condensate in 1995 [1], the special properties of this macroscopic quantum state have been used in a wide range of experiments. One method of utilizing Bose-Einstein condensates is to load them into an optical lattice, which is an array of potentials created through the atoms dipole interaction with laser beams. By utilizing the high controllability of these systems, one can perform quantum simulations of various systems, such as spin chains [34], Dirac cones [37], and artificial gauge fields [6]. Furtermore, the properties of cold atoms in optical lattices is very favourable for experiments in quantum information, where quantum gates can be realised through controlled collisions [14] or Rydberg atoms [31].

This note presents the basic theory of quantum many-body systems in optical lattices.

2 Optical Lattice Potentials

Cold atoms can be trapped in potentials generated by their own dipole interaction with light. Thus, superimposing laser beams allows one to create optical lattices in various shapes and forms. Such dipole traps realised using far-detuned light have important properties, as (i) they are capable of trapping neutral atoms, and (ii) the optical excitation from the trap is very low [11].

Optical lattices are a central component of many experiments, as they not only trap the atoms, but also determines many properties of the system.

2.1 Trapping of Neutral Atoms

Consider a two level atom in the presence of a time-varying electric field $\mathbf{E} = \varepsilon E_0 \exp\{i(\mathbf{k}\mathbf{r} - \omega_L t)\}$, where E_0 is its amplitude, ε is its polarization vector, \mathbf{k} is its wave vector and ω_L is its frequency. The interaction between the atom and the electric field causes a perturbation of the atoms energy levels, otherwise known as the AC Stark-shift. The Hamiltonian describing this interaction is

$$\hat{H}_{int} = \hat{d}\mathbf{E}, \quad (1)$$

where $\hat{d} = -e\hat{r}$ is the electric dipole operator.

Consider equation (1) as a perturbation to the Hamiltonian of the atom. To first order the non-degenerate, time-independent perturbation of the energy of state i reads $E_i^{(1)} = \langle i | \hat{H}_{int} | i \rangle$. However, only states of opposite parity will contribute to the matrix elements of $\langle \hat{H}_{int} \rangle$, as (1) is linear in space. Thus, all first-order perturbation terms cancel.

To second order the perturbation of state i is given by

$$E_i^{(2)} = \sum_{j \neq i} \frac{|\langle j | \hat{H}_{int} | i \rangle|^2}{\varepsilon_i - \varepsilon_j}. \quad (2)$$

Here, the states $|i\rangle$ and $|j\rangle$ and their corresponding energies, ε_i and ε_j , are of the dressed state picture, where one considers the combined system of the atom and the light field [5]. The state $|i\rangle$ represents the atom being in its ground state, while the light field contains n photons. Thus, the energy of the combined state is $\varepsilon_i = n\hbar\omega_L$, when setting the energy of the atomic ground state to zero. Meanwhile, state $|j\rangle$ is when the atom has been excited by absorbing one of the photons of the field. Hence, the energy of this state is $\varepsilon_j = \hbar\omega_0 + (n-1)\hbar\omega_L$. Defining the detuning $\Delta = \omega_L - \omega_0$ allows writing the perturbation in the form

$$E_{g/e}^{(2)} = \pm \frac{|\langle e | \hat{d} | g \rangle|^2}{\Delta} |E_0|^2, \quad (3)$$

where the upper sign is assigned to the atomic ground state $|g\rangle$. This can be rewritten in order to reflect properties of the atom and the field, by considering the intensity of the light, $I = \frac{1}{2}\epsilon_0 c |\mathbf{E}|^2$, and the decay rate of the atom, Γ . Thus, equation (3) can be written as [11]

$$E_{e/g}^{(2)} = \pm \frac{3\pi c^2}{2\omega_0} \frac{\Gamma}{\Delta} I \quad (4)$$

This is the AC-Stark shift, which constitutes the dipole potential. For red detuning ($\Delta < 0$) the ground state will experience a negative shift leading to an attractive potential with depth depending on the intensity of the laser. Similarly, a blue-detuned laser ($\Delta > 0$) will repel the atom. An illustration of this can be seen in figure 1. Since the sign is reversed for the excited state, it is important that the atom remains in the ground state. Thus, one has to minimize the scattering with the optical potential. The scattering rate is given as [11]

$$\Gamma_{sc} = \frac{3\pi c^2}{2\hbar\omega_0^3} \left(\frac{\Gamma}{\Delta} \right)^2 I. \quad (5)$$

As the detuning becomes small, the laser becomes resonant with the atom causing a large increase in scattered photons. Therefore, one has to choose a large detuning in order for the potential to remain

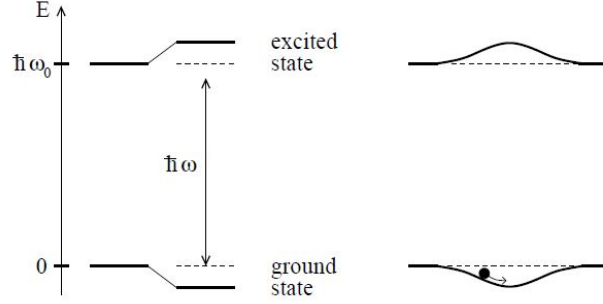


Figure 1: *Light shifts of a two-level atom. Left-hand side, red-detuned light ($\Delta < 0$) shifts the ground state down and the excited state up by same amounts. Right-hand side, a spatially inhomogeneous field like a Gaussian laser beam produces a ground-state potential well, in which an atom can be trapped. Figure and caption are adopted from [11].*

conservative. However, this comes at the cost of a weaker potential. To compensate this, a high laser intensity must be used, in order for the potential to reach sufficient depth. In practise there will be a limit to the laser power available, however, for most alkali-metal atoms the detuning is typically chosen to be large compared to the excited-state hyperfine structure splitting, which provides enough depth while being sufficient to suppress scattering events [3].

2.2 Optical Lattices

The dipole potential in equation (4) scales with the intensity of the laser. Thus, superimposing laser beams allows for creating a multitude of different potentials through the interference patterns of the lasers. A simple standing wave from two counter-propagating light fields will lead to an array of potential wells

$$V(z) = -V_0 \cos^2 kz , \quad (6)$$

where $V_0 = |\frac{3\pi c^2}{2\omega_0^3} \frac{\Gamma}{\Delta} 4I_0|$ from equation (4). In practise, a one dimensional lattice like that of equation (6) is created by shining a single laser beam at a mirror, whereby it interferes with itself. Adding another laser beam in a different direction creates a periodic two dimensional potential. For orthogonal polarization of the two lasers, the resulting potential is purely the sum of the sinusoidal standing wave potential, as no interference term is present [24]. Note, that the lattice is only well defined for distances much smaller than the waist of the laser beams, as the lattice is only present within the overlap of the two beams. Various shapes of the lattice can be achieved by adjusting the angle between the beams, however, the most common setup is using two orthogonal beam creating a square lattice of one dimensional tubes, as seen in figure 2. In order to create a three dimensional lattice, as seen in figure 2, an additional third perpendicular laser beam is needed. In the center of the trap, the lattice potential is then given by

$$V(x, y, z) = -V_0 (\cos^2 kx + \cos^2 ky + \cos^2 kz) , \quad (7)$$

for distances much smaller than the beam waist. In addition to the lattice, an external harmonic confinement will be present due to the Gaussian profile of the laser beams [3].

2.3 Band Structure

Consider a periodic potential as described by equation (6). *Bloch's Theorem* states that energy eigenstates of a periodic potential with lattice vector \mathbf{R} and quasi-momentum q can be written as Bloch waves, which takes the form

$$\phi_{\mathbf{q}}^{(n)}(\mathbf{r}) = e^{i\mathbf{q}\mathbf{r}} u^{(n)}(\mathbf{r}) , \quad (8)$$

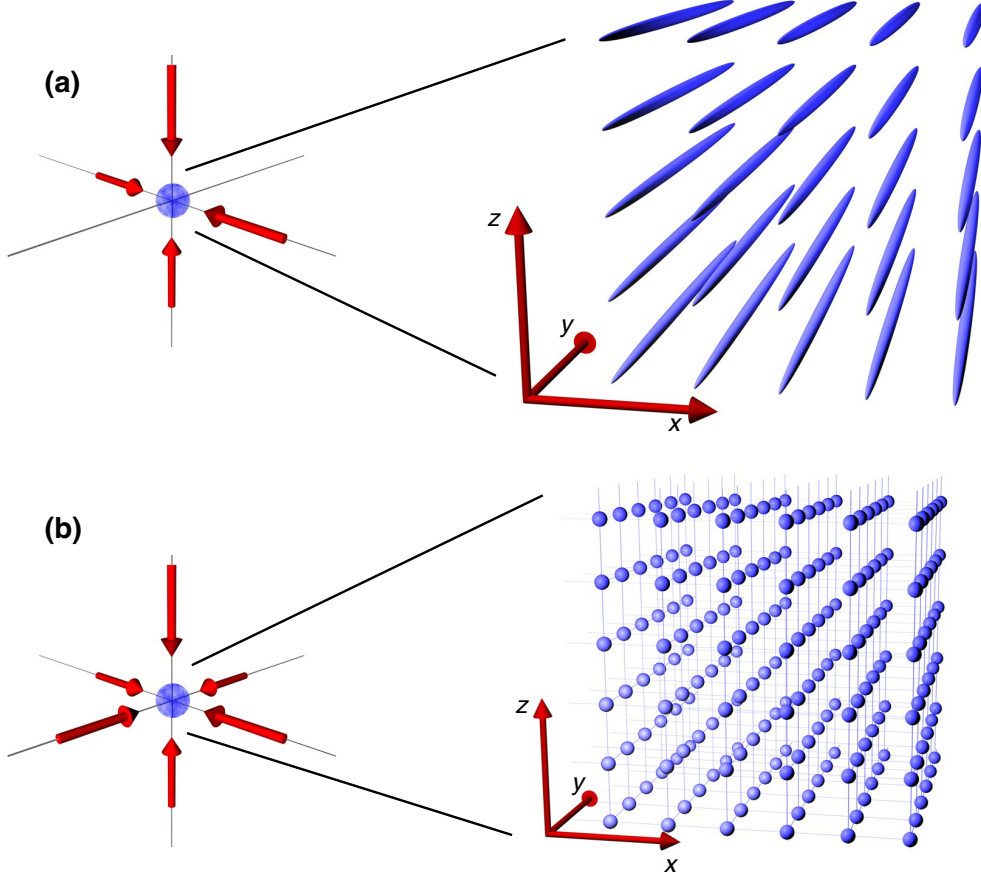


Figure 2: (a) Two dimensional optical lattice formed by two mutually orthogonal laser beams. These tubes have a characteristic cigar shape, due to the Gaussian profile of the lasers. (b) Upon using three orthogonal laser beams, the result is a three dimensional lattice reminiscent of a cubic crystal. Figure is adopted from [43].

which is a plane wave modulated by a function with the same periodicity as the potential $u^{(n)}(\mathbf{r}) = u^{(n)}(\mathbf{r} + \mathbf{R})$. Furthermore, the Bloch waves are periodic in reciprocal space, such that $\psi_{\mathbf{q}}^{(n)}(\mathbf{r}) = \phi_{\mathbf{q} + \mathbf{G}}^{(n)}(\mathbf{r})$, where \mathbf{G} is a reciprocal lattice vector. [18]

This leads to an energy spectrum in the shape of bands with the periodicity of the first *Brillouin Zone*. Bands are denoted by the band index n , and their shape is determined by both the shape and the depth of the potential. The potential depth is often denoted in units of the recoil energy $E_r = \frac{\hbar^2 k^2}{2m}$, where m is the mass of the atom, and k is the photon wave number of the light forming the optical lattice. For $V_0 = 0$, the particles are free, hence the bands will be parabolic. Meanwhile, for $V_0 \rightarrow \infty$, no interaction between different wells of the lattice can take place, as the wavefunctions of the trapped atoms will be confined to their respective well. Thus, the lattice is reduced to an array of independent harmonic oscillators, whereby the bands will appear flat with equal spacing [10].

2.4 Localized States

For lattice potential depths within $5E_r \leq V_0 \leq 8E_r$, the lattice is in the *tight binding limit*. Within this range, the wavefunctions of the trapped atoms will only overlap with other wells in their closest proximity. Thus, interactions between wells are almost purely of nearest neighbour nature. Due to how well localized the wavefunctions are, a basis of Wannier functions is ideal for describing the system.

Wannier functions are related to Bloch functions through the Fourier transform [19]

$$w^{(n)}(\mathbf{r}) = \frac{1}{\sqrt{N_L}} \sum_{\mathbf{q}} e^{-i\mathbf{q}\mathbf{R}} \phi_{\mathbf{q}}^{(n)}(\mathbf{r}) , \quad (9)$$

where N_L is the number of primitive cells of the lattice. The Wannier functions are well localized and centred around the lattice at site \mathbf{R} . In the case of a separable periodic potential, like that of equation (7), the single-particle problem becomes dimensional [20]. Lastly, Wannier functions obey the orthonormality relation

$$\int d^3r \ w^{(n)*}(\mathbf{r} - \mathbf{R}) w^{(n')}(\mathbf{r} - \mathbf{R}') = \delta_{n,n'} \delta_{\mathbf{R},\mathbf{R}'} , \quad (10)$$

thus forming a complete basis [3]. Figure 3 shows Wannier functions plotted for lattice potential depths

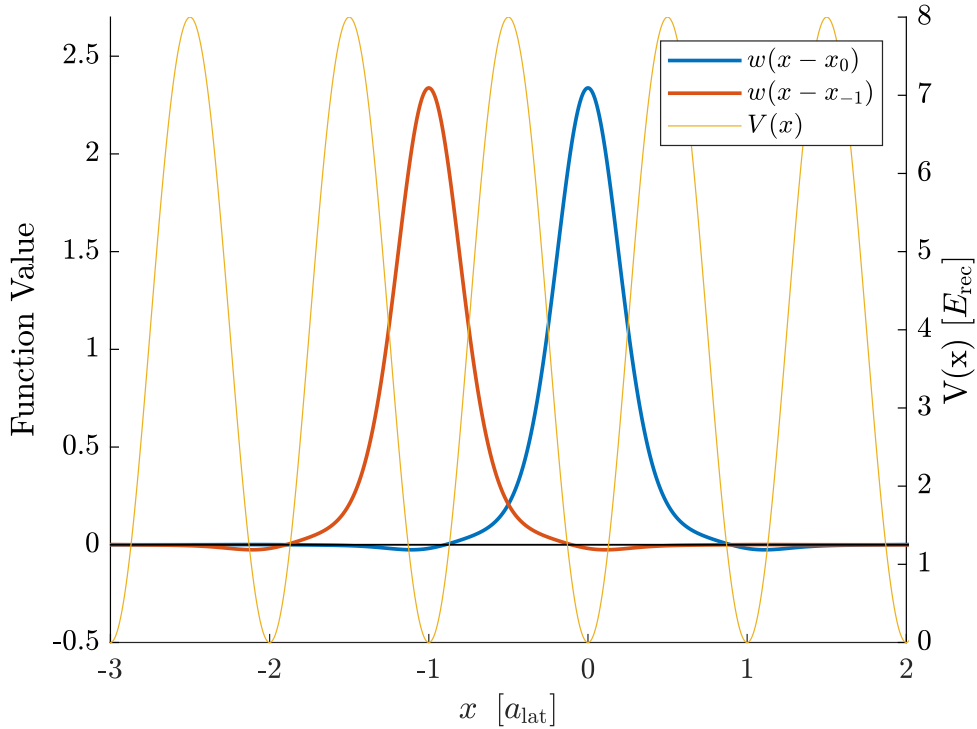


Figure 3: Two one-dimensional Wannier functions plotted for a lattice potential depth of $V_0 = 8E_r$.

within the tight binding limit. As evident from the plot, the functions overlap with only their nearest neighbours. In the case of a more shallow lattice, the functions would extend to further wells, while they will tend towards a Gaussian shape as the lattice depth increases [10].

3 Bose-Einstein Condensates

Bosons are particles of integer spin, whose statistics obey those of a Bose-Einstein distribution

$$n_i = \frac{g_i}{\exp((\varepsilon_i - \mu)/k_B T) - 1} , \quad (11)$$

where i denotes the state, n_i is the population of the state, g_i is its degeneracy, ε_i is its energy, μ is the chemical potential, k_B is the Boltzmann constant, and T is the temperature. One important feature of bosons is that, unlike fermions, multiple particles can occupy the same quantum state. At higher temperatures the energy spectrum is practically continuous, whereby this property has little effect, as two particles occupying the same single-particle state is highly unlikely. However, at low temperatures the energy spectrum systems often become increasingly discrete, hence the statistics of the particles

becomes very important. As evident from equation (11), the population of the ground state diverges as $T \rightarrow 0$. However, even below the finite temperature, T_c , one will observe a macroscopic population of the ground state. T_c is called the critical temperature, and marks the point where multiple particles will start forming a Bose-Einstein Condensate (BEC). [27]

3.1 Non-Interacting Particles

In the case of non-interacting particles and zero temperature, all particles of a Bose gas can be described by identical single-particles wavefunctions $\phi(\mathbf{r}_i)$. Hence, the many-body wavefunction is simply given by the product

$$\Psi(\mathbf{r}_1, \dots, \mathbf{r}_N) = \prod_i^N \phi(\mathbf{r}_i) . \quad (12)$$

Such a product state can be described by a single macroscopic wavefunction

$$\psi(\mathbf{r}) = \sqrt{N} \phi(\mathbf{r}) , \quad (13)$$

where $\phi(\mathbf{r})$ is the wave function of the single-particle state, in which the bosons condensate into [26].

Second-Quantization

When describing Bose-Einstein condensates it is very convenient to work in second quantization, which describes the number of particle in each state rather than the state of each particle.

First, consider a basis of single particle states $\{|n\rangle\}$, namely a Fock basis. In this space particles are created or annihilated through their respective operators

$$\hat{a}_\mu^\dagger |0_\mu\rangle = |1_\mu\rangle . \quad (14)$$

For bosons the creation and annihilation operators fulfill the commutation relations

$$[\hat{a}_\nu, \hat{a}_\mu] = [\hat{a}_\nu^\dagger, \hat{a}_\mu^\dagger] = 0 \quad , \quad [\hat{a}_\nu, \hat{a}_\mu^\dagger] = \delta_{\nu,\mu} , \quad (15)$$

with the number operator given as

$$\hat{n}_\mu = \hat{a}_\mu^\dagger \hat{a}_\mu . \quad (16)$$

In second quantization many-body states are described by the occupation of the individual Fock states. Thus, a creation operator will raise the number of particles in its corresponding state by one, while the annihilation operator will lower it:

$$\hat{a}^\dagger |N_0, N_1, \dots, N_\nu, \dots\rangle = \sqrt{N_\nu + 1} |N_0, N_1, \dots, N_\nu + 1, \dots\rangle \quad (17)$$

$$\hat{a} |N_0, N_1, \dots, N_\nu, \dots\rangle = \sqrt{N_\nu} |N_0, N_1, \dots, N_\nu - 1, \dots\rangle . \quad (18)$$

Likewise, the number operator \hat{n}_μ will count the number of particles in its corresponding state.

These operators can be combined with an orthonormal basis of spatial wavefunctions $\{\phi_k\}$ in order to create field operators

$$\hat{\psi}(\mathbf{r}) = \sum_k \phi_k \hat{a}_k \quad , \quad \hat{\psi}^\dagger(\mathbf{r}) = \sum_k \phi_k^* \hat{a}_k^\dagger , \quad (19)$$

where $\hat{\psi}(\mathbf{r})$ will create a particle at location \mathbf{r} . For bosons the field operators fulfil the commutation relations [4]

$$[\hat{\psi}(\mathbf{r}) , \hat{\psi}^\dagger(\mathbf{r}')] = \delta(\mathbf{r} - \mathbf{r}') \quad , \quad [\hat{\psi}(\mathbf{r}) , \hat{\psi}(\mathbf{r}')] = 0 . \quad (20)$$

Now, consider a gas of non-interacting bosons described by equation (13), where ϵ_k is the energy of the k 'th single-particle state. Due to the completeness of the basis of single-particle wavefunctions, $\{\phi_k\}$, the creation operator can be expressed as

$$\hat{a}_k = \int d^3r \phi_k^*(\mathbf{r}) \hat{\psi}(\mathbf{r}) . \quad (21)$$

Using this, the Hamiltonian can be written as

$$\begin{aligned}
\hat{H}^{(0)} &= \sum_k \epsilon_k \hat{a}_k^\dagger \hat{a}_k \\
&= \sum_k \int d^3 r_1 d^3 r_2 \epsilon_k \phi_k(\mathbf{r}_1) \phi_k^*(\mathbf{r}_2) \hat{\psi}^\dagger(\mathbf{r}_1) \hat{\psi}(\mathbf{r}_2) \\
&= \int d^3 r \hat{\psi}^\dagger(\mathbf{r}) \left(-\frac{\hbar^2}{2m} \nabla^2 + U(\mathbf{r}) \right) \hat{\psi}(\mathbf{r})
\end{aligned} \tag{22}$$

3.2 Weakly Interacting Particles

A characteristic of a BEC is its low temperature and density. Thus, is it a valid approximation to only consider two-particle interactions

$$\hat{H}^{(2)} = \frac{1}{2} \sum_{i \neq j} V(\mathbf{r}_i - \mathbf{r}_j) . \tag{23}$$

At low energies all interactions can be considered of s-wave nature, because all waves of higher angular momentum are reflected by the centrifugal barrier. Furthermore, for cold gases the thermal de Broglie wavelength is much larger than the effective extension of the interaction potential. Therefore, the actual shape of the scattering potential is irrelevant, hence one can replace it with a pseudo-potential

$$V(\mathbf{r} - \mathbf{r}') = g \delta(\mathbf{r} - \mathbf{r}') = \frac{4\pi\hbar^2 a}{m} \delta(\mathbf{r} - \mathbf{r}') , \tag{24}$$

which results in the same scattering phase as the real, more complicated scattering potential. Thus, the interaction of cold atoms is fully determined by the scattering length, a [10].

Introducing the density operator

$$\hat{\rho}(\mathbf{r}) = \hat{\psi}^\dagger(\mathbf{r}) \hat{\psi}(\mathbf{r}) , \tag{25}$$

allows writing $\hat{H}^{(2)}$ in second quantization

$$\hat{H}^{(2)} = \frac{1}{2} \int d^3 r_1 d^3 r_2 V(\mathbf{r}_1 - \mathbf{r}_2) \hat{\psi}^\dagger(\mathbf{r}_1) \hat{\psi}(\mathbf{r}_1) \left(\hat{\psi}^\dagger(\mathbf{r}_2) \hat{\psi}(\mathbf{r}_2) - \delta(\mathbf{r}_1 - \mathbf{r}_2) \right) \tag{26}$$

$$= \frac{1}{2} \int d^3 r_1 d^3 r_2 \hat{\psi}^\dagger(\mathbf{r}_1) \hat{\psi}^\dagger(\mathbf{r}_2) V(\mathbf{r}_1 - \mathbf{r}_2) \hat{\psi}(\mathbf{r}_1) \hat{\psi}(\mathbf{r}_2) . \tag{27}$$

Combining this with the basic Hamiltonian of equation (22), gives the full Hamiltonian in second quantization

$$\hat{H} = \hat{H}^{(0)} + \hat{H}^{(2)} \tag{28}$$

$$= \int d^3 r \hat{\psi}^\dagger(\mathbf{r}) \left(-\frac{\hbar^2}{2m} \nabla^2 + U(\mathbf{r}) \right) \hat{\psi}(\mathbf{r}) + \frac{1}{2} \int d^3 r_1 d^3 r_2 \hat{\psi}^\dagger(\mathbf{r}_1) \hat{\psi}^\dagger(\mathbf{r}_2) V(\mathbf{r}_1 - \mathbf{r}_2) \hat{\psi}(\mathbf{r}_1) \hat{\psi}(\mathbf{r}_2) \tag{29}$$

Using this Hamiltonian to try and solve the Heisenberg equations of motion for the field operators leads to

$$i\hbar \frac{\partial}{\partial t} \hat{\psi}(\mathbf{r}) = \left[\hat{\psi}(\mathbf{r}) , \hat{H} \right] = \left(-\frac{\hbar^2}{2m} \nabla^2 + U(\mathbf{r}) + g \hat{\psi}^\dagger(\mathbf{r}) \hat{\psi}(\mathbf{r}) \right) \hat{\psi}(\mathbf{r}) , \tag{30}$$

which is not solvable in general. However, in the scenario of a BEC the scattering length a is much less than the mean inter-particle distance, such that $na^3 \ll 1$, where n is the density of the gas. In this regime the mean-field approximation is viable

$$\hat{\psi}(\mathbf{r}) = \psi(\mathbf{r}) + \delta\hat{\psi}(\mathbf{r}) , \tag{31}$$

where $\psi(\mathbf{r})$ is the mean-field given by equation (13), and $\delta\hat{\psi}(\mathbf{r})$ is fluctuations from the mean. If $\langle \delta\hat{\psi}(\mathbf{r}) \rangle = 0$, the fluctuations can be neglected, leading to the Gross-Pitaevskii equation [12, 28]

$$i\hbar \frac{\partial}{\partial t} \hat{\psi}(\mathbf{r}) = \left(-\frac{\hbar^2}{2m} \nabla^2 + U(\mathbf{r}) + g|\psi(\mathbf{r})|^2 \right) \psi(\mathbf{r}) . \tag{32}$$

The Gross-Pitaevskii equation is very similar to the Schrödinger equation with exception of the non-linear term $g|\psi(\mathbf{r})|^2$, which can make the equation hard to solve in regions of low density.

4 Bose-Hubbard Model of Interacting Bosons in a Lattice

The Bose-Hubbard model describes weakly interacting bosons in a periodic lattice within the tight binding limit. The model is perfectly capable of predicting results with good accuracy, however, two conditions must hold in order for it to be valid: (i) both the thermal energy and the mean interaction energy at a single site must be much smaller than the separation to first excited band, $\hbar\omega_0$, and (ii) the Wannier functions decay essentially within the length of the lattice constant [3].

Under these conditions one is assured that only the lowest band is taken into account, and that only nearest-neighbour interactions take place.

The Bose-Hubbard model is interesting, as it supports two distinct phases: the Superfluid phase and the Mott-Insulator phase. Furthermore, the model contains effects such as a quantum phase transition between the two phases mentioned above, which can be crossed without any change in external parameters.

4.1 The Bose-Hubbard Hamiltonian

Consider the Hamiltonian for bosonic particles in a trapping potential (described by equation (29)) in one dimension. For a periodic lattice potential in the tight binding limit it is favourable to work in a basis of localized Wannier functions. Expanding the field operators of the Hamiltonian in equation (29) in the Wannier basis yields [15]

$$\begin{aligned} \hat{H} = & \int dx \sum_{ij} w^*(x - x_i) \hat{a}_i^\dagger \left(-\frac{\hbar^2}{2m} \nabla^2 + V(x) \right) w(x - x_j) \hat{a}_j \\ & + g \int dx \sum_{ijkl} w^*(x - x_i) w^*(x - x_j) w(x - x_k) w(x - x_l) \hat{a}_i^\dagger \hat{a}_j^\dagger \hat{a}_k \hat{a}_l \end{aligned} \quad (33)$$

$$= - \sum_{ij} J_{ij} \hat{a}_i^\dagger \hat{a}_j + \sum_{ijkl} U_{ijkl} \hat{a}_i^\dagger \hat{a}_j^\dagger \hat{a}_k \hat{a}_l, \quad (34)$$

where

$$J_{ij} = - \int dx w^*(x - x_i) \left(-\frac{\hbar^2}{2m} \nabla^2 + V(x) \right) w(x - x_j) \quad (35)$$

$$U_{ijkl} = g \int dx w^*(x - x_i) w^*(x - x_j) w(x - x_k) w(x - x_l) \quad (36)$$

Since the system is periodic, one can consider a single site, $i = 0$, as representative for the entire lattice. In this way, the different terms of the Hamiltonian can be interpreted as follows:

- J_{00} = constant energy offset
- J_{01} = "overlap matrix element" to neighbouring site
- $J_{02-0\infty}$ = "overlap matrix element" to further sites
- U_{0000} = on-site interaction for two particles
- U_{0ii0} = interaction off-site
- U_{0001} = interaction + tunnelling, off-site

Due to the rapid decay of the Wannier functions, the overlap of wavefunctions is limited to their nearest neighbour. Dropping all exponentially suppressed terms and constant offsets yields the Bose-Hubbard Hamiltonian

$$\hat{H} = -J \sum_{\langle i,j \rangle} \hat{a}_i^\dagger \hat{a}_j + \frac{U}{2} \sum_i \hat{n}_i (\hat{n}_i - 1) + \sum_i \varepsilon_i \hat{n}_i, \quad (37)$$

where $J = J_{01}$, $\langle i, j \rangle$ only counts neighbouring pairs, and

$$U = U_{0000} = g \int dx |w(x)|^4. \quad (38)$$

The first term of the Bose-Hubbard Hamiltonian describes the kinematics within the model, which takes the form of tunneling between neighbouring sites. This is interpreted by annihilating a particle at site j while creating a particle at site i . The second term describes the interaction between particles within a single site, and the last term $\sum_i \varepsilon_i \hat{n}_i$ takes into account a possible potential offset at different sites.

4.2 Phases of the Bose-Hubbard Model

The Bose-Hubbard model supports two quantum phases: The *Superfluid* phase and the *Mott Insulator* phase. These phases depend on the ratio J/U , and can be described separately by examining the ground state of the Bose-Hubbard Hamiltonian in the two extreme limits of the J/U ratio.

Superfluid Phase

For a system where the tunneling matrix element J is dominant, the lowest energy is obtained by delocalizing the atoms over the entire lattice. Thereby the wavefunction will be a product over single-particle states, and the system will be a superfluid [10].

Consider the case of negligible interactions and a lattice of equal depth. In this scenario the Bose-Hubbard Hamiltonian reduces to

$$\hat{H} = \hat{H}_J = -J \sum_{\langle i, j \rangle} \hat{a}_i^\dagger \hat{a}_j, \quad (39)$$

which is completely periodic within the lattice due to the lack of site specific terms. This leads to solutions in the shape of Bloch waves. The Fourier transform of the annihilation and creation operators

$$\hat{a}_j = \frac{1}{N_L} \sum_q e^{iqx_j} \hat{a}_q, \quad \hat{a}_j^\dagger = \frac{1}{N_L} \sum_q e^{-iqx_j} \hat{a}_q^\dagger \quad (40)$$

allows for writing the Hamiltonian (39) in momentum space

$$\hat{H}_J = -J \sum_{q=-\infty}^{\infty} \left(e^{-iqd} + e^{ikd} \right) \hat{n}_q, \quad (41)$$

where d is the lattice distance. In momentum space the Hamiltonian is diagonal and results in a continuous energy spectrum

$$E_q = -2J \cos(qd). \quad (42)$$

Thus, the excitation spectrum of the Superfluid is said to be gapless.

The lowest energy of the system is obtained for $q = 0$, whereby the ground state is

$$|\Psi_{SF}\rangle = \frac{1}{\sqrt{N!}} \left(\hat{a}_{q=0}^\dagger \right)^N |0\rangle = \frac{1}{\sqrt{N!}} \left(\frac{1}{N_L} \sum_{j=1}^{N_L} \hat{a}_j^\dagger \right)^N |0\rangle, \quad (43)$$

where N is the number of particles. This state supports a well defined macroscopic phase on each lattice site, since the many-body state is a product over identical single-particle states [10]. With all particles condensed into the $q = 0$ momentum space state, the particles are completely de-localized in real space, which can be seen by taking the Fourier transform. For large N , $N_L \rightarrow \infty$ at fixed density N/N_L the state becomes indistinguishable from having coherent states, $|\alpha\rangle$, on all lattice sites [3]

$$|\Psi_{SF}\rangle \approx \prod_j \left(e^{\sqrt{N/N_L} \hat{a}_j^\dagger} \right) |0\rangle = |\alpha_1\rangle \otimes |\alpha_2\rangle \otimes \dots \quad (44)$$

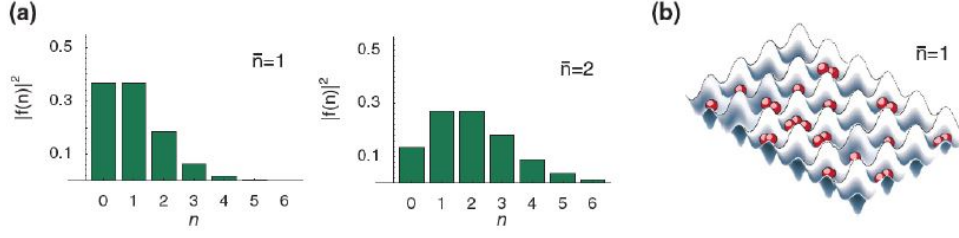


Figure 4: (a) The statistics for the number of particles per lattice site n for a filling fraction of $\bar{n} = 1$ and $\bar{n} = 2$ in the superfluid phase. (b) Illustration of the particles in the lattice. There will be large fluctuations of the number of particles found at each lattice site at a given time. Figure and caption adopted from [10].

Since bosonic operators at different sites commute, the superfluid state can be factorized into a product of local coherent states. Coherent states are eigenstates of the annihilation operator $\hat{a}|\alpha\rangle = \alpha|\alpha\rangle$, where $|\alpha|^2$ can be considered the particle density of the system, and $\alpha = |\alpha|e^{i\phi}$, with ϕ being a global phase. Utilizing the coherent state form of the wavefunction (eq. (44)) the average filling fraction \bar{n} can be calculated

$$\bar{n} = \langle \hat{n}_i \rangle = \langle \Psi_{SF} | \hat{a}_j^\dagger \hat{a}_j | \Psi_{SF} \rangle = \frac{N}{N_L}, \quad (45)$$

as well as the fluctuations of particle number per site

$$\frac{\sqrt{\Delta \bar{n}^2}}{\bar{n}} \sim \frac{1}{\sqrt{\bar{n}}}. \quad (46)$$

Therefore, the probability distribution for the number of atoms at a given site is Poissonian, which can be seen in figure 4. Due to the relatively large fluctuation of particle number per site, one would find a somewhat random number of atoms at a given site in a measurement. In the presence of a finite interaction, U , the resulting distribution would be sub Poissonian due to number squeezing [10].

Mott-Insulator Phase

In the case of negligible tunneling, only the on-site interaction between the atoms has to be taken into account, and the Bose-Hubbard Hamiltonian reduces to

$$\hat{H} = \hat{H}_U = \frac{U}{2} \sum_i \hat{n}_i (\hat{n}_i - 1), \quad (47)$$

which is quadratic in \hat{n}_i . This heavily penalizes having multiple particles at the same site. Thus, the ground state of the system will be an equal distribution of all the particles throughout the lattice. Any fluctuations from this average will increase the energy, whereby the phase is said to be incompressible [9]. This incompressibility can be formulated as $\frac{\partial n}{\partial \mu} = 0$, where μ is the chemical potential, and is the defining property of the Mott-Insulator [3].

Consider the case $\bar{n} = 1$, (average filling of one particle per site). This can be described by a state with a single particle located on each site [3]

$$|\Psi_{Mott}\rangle = \prod_j \hat{a}_j^\dagger |0\rangle. \quad (48)$$

This is a simple product of local Fock states with precisely one atom per site, which is illustrated in figure 5. This is the configuration of atoms, which minimizes the energy with regards to the Hamiltonian of eq. (47). Any fluctuation from unit occupancy will increase the energy, as merely a single double occupancy will increase the energy by U . Thereby, unlike the energy spectrum of the Superfluid, the Mott-Insulator spectrum is gapped. As long as the gain in kinetic energy due to hopping, J , is smaller than the on-site interaction, U , the atoms remain localized. However, for $J > 0$ the ground state is no longer a the simple product state described in eq. (48) [3]. As J increases, the

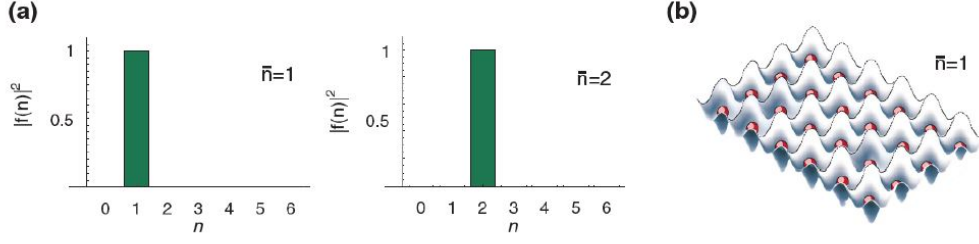


Figure 5: (a) The statistics for the number of particles per lattice site n in the Mott-insulator phase, for a filling fraction of $\bar{n} = 1$ and $\bar{n} = 2$. (b) Illustration of the particles in the lattice. The particles do not hop around in the lattice and are equally distributed. Caption and figure are adapted from [10].

gap in the excitation spectrum will gradually decrease until the transition to the superfluid is reached and the spectrum becomes gapless.

While the Mott-Insulator phase has complete localization, the phases on the individual sites have obtained maximum uncertainty. Therefore, no phase coherence between different sites is present [10].

4.3 Phase Transition

Consider once again the full Bose-Hubbard Hamiltonian (37). The superfluid and the Mott-Insulator account for the two limits of the ratio J/U , however, in order to understand the full phase diagram of the Bose-Hubbard model one has to derive the U_{crit} for which the phase transition occurs. There are several ways of doing this - one of them is looking at a mean-field solution of the Bose-Hubbard model. For simplicity the scenario $T = 0$ is treated here. Even without a change of temperature the phase transition still happens, hence the label *quantum phase transition* referring to the fact that the phase transition can happen without change of external parameters [30].

Applying the mean field approximation to the annihilation operator yields

$$\hat{a}_j = \psi + \delta\hat{a}_j, \quad (49)$$

where ψ is the locally constant mean field, and $\delta\hat{a}_j$ is the fluctuation term. Inserting this into the Hamiltonian yields

$$\hat{H}_{MF} = -J \sum_{\langle i,j \rangle} \left(\psi^* + \delta\hat{a}_i^\dagger \right) (\psi + \delta\hat{a}_j) + \text{int.} \quad (50)$$

Instead of considering the Hamiltonian as a whole, it can be considered as a sum of local on-site Hamiltonians

$$\hat{H}_{MF} = \sum_i \hat{h}_i. \quad (51)$$

For a homogeneous system $\hat{h}_i = \hat{h}_j \ \forall i, j$. One can write the kinetic part of the Hamiltonian locally if one assumes small fluctuations from the mean field

$$\begin{aligned} \hat{a}_i^\dagger \hat{a}_j &= \left(\psi^* + \delta\hat{a}_i^\dagger \right) (\psi + \delta\hat{a}_j) \\ &= \psi^* \psi + \psi^* \delta\hat{a}_j + \psi \delta\hat{a}_i^\dagger + \delta\hat{a}_i^\dagger \delta\hat{a}_j \\ &\approx \psi^* \psi + \psi^* (\hat{a}_j - \psi) + \psi (\hat{a}_i^\dagger - \psi^*) \\ &= \psi^* \hat{a}_j + \psi \hat{a}_i^\dagger - \psi^* \psi \end{aligned} \quad (52)$$

Since every term only contains a single index, one can let $j \rightarrow i$ leaving a local description

$$\hat{h}_i = Jz\psi^*\psi - Jz \left(\psi^* \hat{a}_i + \psi \hat{a}_i^\dagger \right) + \frac{U}{2} \hat{n}_i (\hat{n}_i - 1) + \mu_i \hat{n}_i, \quad (53)$$

where z is the number of neighbours of site i [38]. The last term added is a chemical potential, which is needed due to the lack of a fixed particle number locally prompting the use of a Grand Canonical

Ensemble description.

Landau theory is a general theory regarding phase transitions, which states that in the vicinity of a critical point, one may expand the free energy in a power series of some order parameter m . Thus, in order to find U_{crit} , equation (53) must be minimized. In this case this order parameter is the mean field, hence

$$E_{MF} = \text{const.} + a|\psi|^2 + b|\psi|^4 + \dots \quad (54)$$

Plotting this yields an effective potential, seen in figure 6, which dictates certain properties of the system.

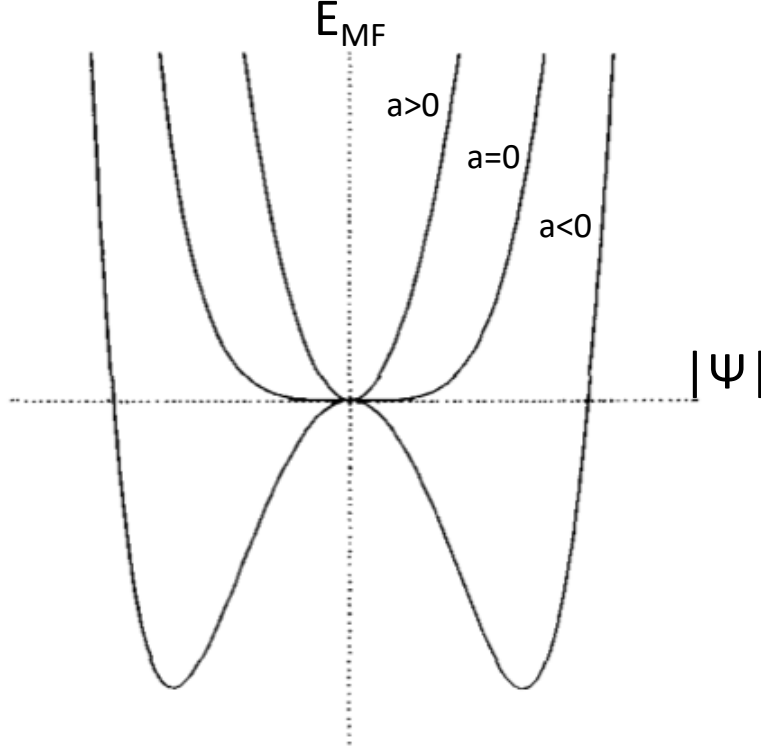


Figure 6: Effective potential as function of the order parameter ψ . For $a < 0$ the properties of the potential change from having a single, central minimum to having two separated minima. This is an example of spontaneous symmetry breaking associated with a phase transition. [29]

The potential is symmetric in the complex plane allowing the system to have any phase. Furthermore, at some critical value of a , (here $a = 0$), the potential will shift from having a single central minimum to having minima at some $|\psi|$. This is a U(1) spontaneous symmetry breaking, which in Landau theory is associated with a phase transition [29]. Thus, one needs to compute the parameter a of equation (54). One method is using second order perturbation theory, where the $\hat{\psi} = 0$ case is solved exactly, followed by adding a small $\hat{\psi}$ as a perturbation

$$\hat{h}_i^{(0)} = Jz(\psi^*\psi) + \frac{U}{2}\hat{n}_i(\hat{n}_i - 1) - \mu\hat{n}_i. \quad (55)$$

Since the zero-order solution only contains number operators, the ground state of the solution can be expressed in the Fock basis:

$$\begin{aligned} |0\rangle & \text{ for } \mu < 0 \\ |1\rangle & \text{ for } 0 \leq \mu < U \\ |2\rangle & \text{ for } U \leq \mu < 2U \\ & \vdots \end{aligned}$$

Adding on the first order perturbation to the energy

$$E_i^{(1)} = \langle n | \delta \hat{a}_i | n \rangle = 0 , \quad (56)$$

yields nothing, thus requiring the use of a second order perturbation

$$E_i^{(2)} = \sum_{n \neq g} \frac{|\langle n | \delta \hat{h}_i | g \rangle|^2}{E_g^{(0)} - E_n^{(0)}} . \quad (57)$$

Here the perturbation Hamiltonian

$$\hat{h}_i = -zJ \left(\hat{a}_i \psi^* + \psi \hat{a}_i^\dagger \right) \quad (58)$$

contains only single creation/annihilation operators, whereby

$$\langle n | \delta \hat{h}_i | g \rangle = 0 \quad \text{for} \quad |n - g| \neq 1 . \quad (59)$$

Hence, for each annihilation and creation operator only two matrix elements will give a contribution

$$\begin{aligned} \langle n | \hat{a}_i^\dagger | n - 1 \rangle &= \sqrt{n} \langle n | n \rangle \\ \langle n + 1 | \hat{a}_i^\dagger | n \rangle &= \sqrt{n + 1} \langle n | n \rangle \\ &\vdots \end{aligned}$$

With this the second order perturbation energy reduces to

$$\begin{aligned} E_i^{(2)} &= (Jz)^2 |\psi|^2 \left(\frac{n}{E_g^{(0)} - E_{n-1}^{(0)}} + \frac{n+1}{E_n^{(0)} - E_{n+1}^{(0)}} \right) \\ &= (Jz)^2 |\psi|^2 \left(\frac{n}{U(n-1) - \mu} + \frac{n+1}{\mu - Un} \right) . \end{aligned} \quad (60)$$

Since a is the pre-factor of all terms proportional $|\psi|^2$, collecting those across from all the perturbations yields the approximation

$$a = Jz + (Jz)^2 |\psi|^2 \left(\frac{n}{U(n-1) - \mu} + \frac{n+1}{\mu - Un} \right) . \quad (61)$$

As stated earlier U_{crit} can be found from $a = 0$:

$$0 \stackrel{!}{=} 1 + \frac{n}{\bar{U}(n-1) - \bar{\mu}} + \frac{n+1}{\bar{\mu} - \bar{U}n} , \quad (62)$$

with $\bar{m}u = \frac{\mu}{Jz}$ and $\bar{U} = \frac{U}{Jz}$. Finally, the solution for the chemical potential is [38]

$$\bar{m}u_{\pm} = \frac{1}{2} \left(\bar{U}(2n-1) \pm \frac{1}{2} \sqrt{\bar{U}^2 - 2\bar{U}(2n+1)} \right) . \quad (63)$$

Understanding this result can be done by examining figure ??, which displays a phase diagram of the Bose-Hubbard model. The μ_{\pm} curves enclose the region where the system is in the Mott Insulating phase. $(1/\bar{U})_{crit} = (J/U)_{crit}$ can be read off the graph from the point, where the two curves μ_{\pm} meet. As mentioned earlier, no fluctuations take place in the Mott Insulator, whereby the particle number per site is well defined, while it for the superfluid can take many values. As the chemical potential increases each site can accommodate more particles as long as the increase in chemical potential compensates the increased energy due to interactions between the particles.

The mean-field solution of the Bose-Hubbard model is only an approximation, which proves quite inaccurate for one dimension. This is seen when comparing the critical ratio for the mean-field approach, $(U/J)_{crit}^{MF,1D} = 11.66$, with numerical results computed using the DMRG method, $(U/J)_{crit}^{DMRG,1D} =$

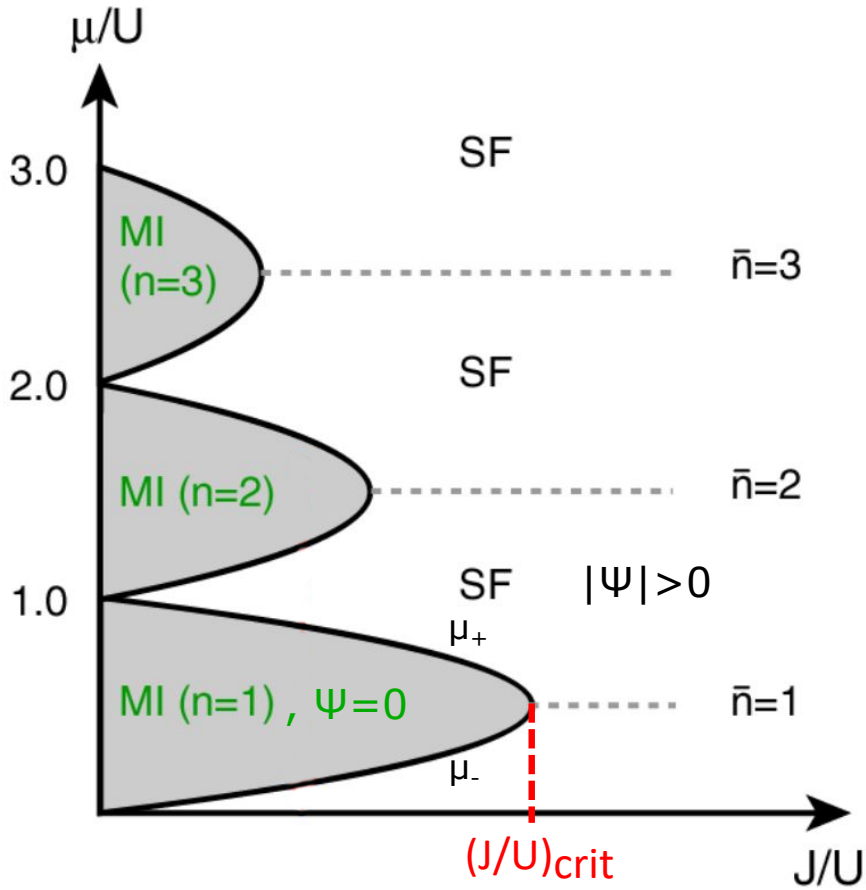


Figure 7: Phase diagram of Bose-Hubbard model for $T = 0$. Grey areas mark the Mott Insulator phase for different number of particles per site, while white regions mark the superfluid phase. [10]

3.37, [22]. Nevertheless, it gives a good intuitive feeling of the physics taking place and how one can describe them without resorting to diagonalizing the Hamiltonian.

5 Matrix Product States

Matrix product states is a parametrization of one dimensional states, which only considers the relevant part of the full Hilbert space, thus greatly simplifying numerical computations within the system. The main idea is to parametrize the coefficients of a general linear combination of states as a product of matrices, with a single matrix for each physical site.

SKRIV INTRO OM MPS MED REFERENCER TIL LITTERATUR - quick "history" * known for a long time * algorithms standardised by White - why do we need MPS description? (copy from numerics chapter) * Do quick analysis of size of Hilbert space * mention efficient application of operators - usage of MPS (mention DMRG and tDMRG here)

5.1 Entanglement in Quantum Systems and Area Laws

Entanglement is a fundamental property of quantum mechanics responsible for correlating different degrees of freedom within a quantum system. Thus, the individual parts of a quantum system can not be described alone, as entanglement with the remainder of the system has to be taken into consideration. This drastically complicates any description of the system, which is why exact descriptions of many-body systems are almost impossible. CITE

The measure of entanglement within a quantum many-body system is the entanglement entropy,

although it is often more instructive to look at the bipartite entanglement entropy, which measures the entanglement between two partitions of the system. Consider a bipartition of the Hilbert space $\mathcal{H} = \mathcal{H}_A \otimes \mathcal{H}_B$. A state $|\psi\rangle \in \mathcal{H}$ can be decomposed using the *Schmidt decomposition* as

$$|\psi\rangle = \sum_{\alpha} \Lambda_{\alpha} |\alpha\rangle_A \otimes |\alpha\rangle_B , \quad (64)$$

where the states $\{|\alpha\rangle_{A(B)}\}$ form an orthonormal basis of $\mathcal{H}_{A(B)}$, and $\Lambda_{\alpha} \geq 0$ are the Schmidt coefficients fulfilling $\sum_{\alpha} \Lambda_{\alpha}^2 = 1$ CITE. If only one term contributes to the Schmidt decomposition, the state is obviously a product state i.e. the two parts of the Hilbert space are not mutually entangled. More terms, on the other hand, implies that the state is entangled. From the Schmidt decomposition one can determine the reduced density matrix

$$\rho_A = \text{Tr}_B |\psi\rangle \langle\psi| = \sum_{\alpha} \Lambda_{\alpha}^2 |\alpha\rangle_A \langle\alpha|_A . \quad (65)$$

From this one can determine the entanglement entropy of the chosen bipartition, which is defined as the von-Neumann entropy, S , of the reduced density matrix given by CITE

$$S = - \sum_{\alpha} \Lambda_{\alpha}^2 \log \Lambda_{\alpha}^2 . \quad (66)$$

Quite often, one is not concerned with the actual entanglement entropy of the system, but rather how it scales when the region in question grows in size. It is natural to assume that the entanglement entropy scales with the systems size, the *volume* of the system, since this is the case for thermal systems. However, ground states of quantum many-body systems often follow an *area law*, meaning that the scaling of the entropy is linear in the boundary area of the region. This is especially the case for systems with a gapped and local Hamiltonian [7]. This observation is the key to the success of numerical computations for one-dimensional systems, since the ground state can be described with relatively few parameters. Thus, one only has to consider a small region of the Hilbert space when performing a variational search for the ground state.

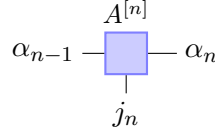
For one dimensional systems the boundry of a bipartition consists of only one site. Hence, following an area law means that the entropy will be bounded by a constant, which is independent of the size of both the system and the subsystem. This can be understood intuitively as only degrees of freedom within the correlation length, ξ , of the boundary will be entangled, no matter where in the system the boundary is present [?].

5.2 Diagrammatic Representation of Matrix Product States

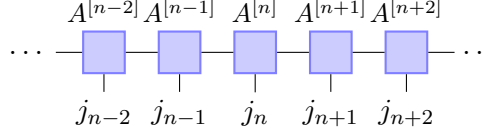
A powerful method for describing one dimensional systems is through *Matrix Product States*, which in principle are a representation of the system generated through a series of Schmidt decompositions. Thus, it immediately follows that MPS's satisfy an area law, making them excellent at describing ground states.

Matrix product states is a parametrization of one dimensional states, which only considers the relevant part of the full Hilbert space, thus greatly simplifying numerical computations within the system. The main idea is to parametrize the coefficients of a general linear combination of states as a product of matrices, with a single matrix for each physical site.

An MPS and any operations on it can be described graphically using tensor networks, which allows for a visual overview of the mathematics taking place. For instance, one can represent a single tensor in the MPS as seen in figure 14i, where horizontal legs are called *bond dimensions* and represent the indices of the local matrix. The vertical legs correspond to the physical index of the site. These tensors can be contracted by connecting the legs of tensors as seen in figure 14ii. Connected lines are summed over just like matrix multiplication.



(i) *Single tensor of an MPS. The horizontal legs are the indices of the local matrix, while the vertical leg is physical index of the site.*



(ii) *Simple connected network of tensors. Connected lines are summed over.*

Singular Value Decomposition (SVD)

A central tool for parametrizing a state as an MPS is the *singular value decomposition*. An arbitrary matrix, A , of dimensions $(M \times N)$ can be decomposed into three matrices as

$$A = USV^\dagger, \quad (67)$$

where each of the matrices contains unique properties [?]:

- U is of dimension $(M \times \min(M, N))$ and has orthonormal columns, meaning $U^\dagger U = I$. If $M \leq N$ then it is also unitary $UU^\dagger = I$.
- S is a positive, diagonal $(\min(M, N) \times \min(M, N))$ matrix. The diagonal elements are singular values, and the number of non-zero entries is the Schmidt rank of A .
- V^\dagger is of dimension $(\min(M, N) \times N)$ and has orthonormal rows, meaning $V^\dagger V = I$. If $M \geq N$ then it is also unitary $VV^\dagger = I$.

These properties are very important when building the MPS. Figure 15 shows how this is displayed using diagrams.

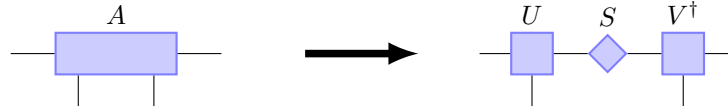


Figure 9: SVD of a tensor A shown diagrammatically.

5.3 Construction of an MPS

Consider a chain of N sites with each site having a d -dimensional local Hilbert space $\{|j_n\rangle\}$, where $n = 1, \dots, N$. There are several ways of constructing an MPS from an arbitrary quantum state of this chain

$$|\psi\rangle = \sum_{j_1, \dots, j_N} c_{j_1 \dots j_N} |j_1, \dots, j_N\rangle. \quad (68)$$

One method is to apply successive SVD's, where the ordering of these operations determines the canonical form of the SVD. The construction of an MPS requires a series of steps:

1. The d^N -dimensional vector $c_{j_1 \dots j_N}$ is reshaped into a $(d \times d^{N-1})$ matrix Ψ , where

$$\Psi_{j_1, (j_2 \dots j_N)} = c_{j_1 \dots j_N}. \quad (69)$$

Performing an SVD on said matrix yields

$$c_{j_1 \dots j_N} = \Psi_{j_1, (j_2 \dots j_N)} = \sum_{\alpha_1}^d U_{j_1, \alpha_1} S_{\alpha_1, \alpha_1} (V^\dagger)_{\alpha_1, (j_2 \dots j_N)} = \sum_{\alpha_1}^d A_{\alpha_1}^{j_1} \Psi_{(\alpha_1 j_2), (j_3 \dots j_N)} , \quad (70)$$

where U has been decomposed into d row vectors A^{j_1} with entries $A_{\alpha_1}^{j_1} = U_{j_1, \alpha_1}$. Furthermore, S and V^\dagger has been multiplied and reshaped into a $(d^2 \times d^{N-2})$ matrix.

2. The new matrix $\Psi_{(\alpha_1 j_2), (j_3 \dots j_N)}$ is subjected to an SVD decomposition

$$c_{j_1 \dots j_N} = \sum_{\alpha_1}^d \sum_{\alpha_2}^{d^2} A_{\alpha_1}^{j_1} U_{(\alpha_1 j_2), \alpha_2} S_{\alpha_2, \alpha_2} (V^\dagger)_{\alpha_2, (j_3 \dots j_N)} = \sum_{\alpha_1}^d \sum_{\alpha_2}^{d^2} A_{\alpha_1}^{j_1} A_{\alpha_1, \alpha_2}^{j_2} \Psi_{(\alpha_2 j_3), (j_4 \dots j_N)} , \quad (71)$$

where U has been decomposed into d matrices A^{j_2} with entries $A_{\alpha_1, \alpha_2}^{j_2} = U_{(\alpha_1 j_2), \alpha_2}$. The matrix $\Psi_{(\alpha_2 j_3), (j_4 \dots j_N)}$ has dimensions $(d^3 \times d^{N-3})$.

3. This pattern of applying and SVD to the matrix, replacing U and reshaping into a new matrix is continued throughout the chain leaving

$$c_{j_1 \dots j_N} = \sum_{\alpha_1, \dots, \alpha_{N-1}} A_{\alpha_1}^{j_1} A_{\alpha_1, \alpha_2}^{j_2} \dots A_{\alpha_{N-2}, \alpha_{N-1}}^{j_{N-1}} A_{\alpha_{N-1}, \alpha_N}^{j_N} = A^{j_1} A^{j_2} \dots A^{j_{N-1}} A^{j_N} , \quad (72)$$

where the sum can be recognized as a matrix multiplication allowing a neater notation. The whole process is illustrated in figure 16.

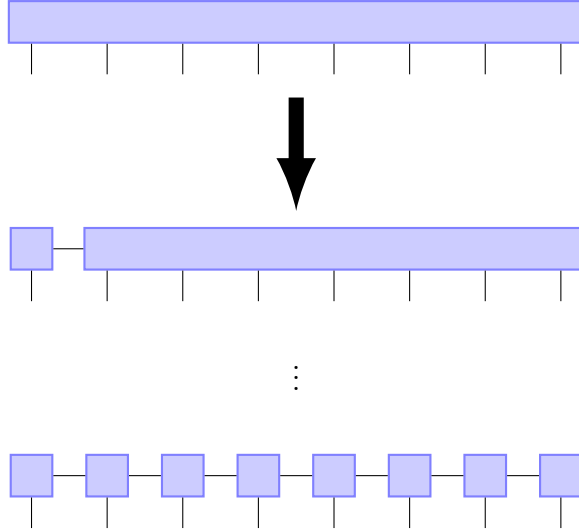


Figure 10: Diagrammatic representation of the construction of a left-canonical MPS from an arbitrary quantum state through successive SVD's.

Thus, one can rewrite the original state (equation 84) as a matrix product state [?]

$$|\psi\rangle = \sum_{j_1, \dots, j_N} A^{j_1} A^{j_2} \dots A^{j_{N-1}} A^{j_N} |j_1, \dots, j_N\rangle . \quad (73)$$

It is worth examining the properties of this representation. Looking at the dimensions of the components of an SVD one can see that the dimensions of the matrices A^{j_n} follow a stair-like structure $(1 \times d), (d \times d^2), \dots, (d^{N/2-1} \times d^{N/2}), (d^{N/2} \times d^{N/2-1}), \dots, (d \times 1)$, where N is taken as even for simplicity. Hence, the matrix dimensions increase exponentially making exact computations practically impossible. However, certain simplifications can be made, greatly increasing computational speed without sacrificing much precision at all. First, one only needs to keep the non-zero singular values of the matrices S , thus

reducing the dimensional factor from d to r_n , where $r_n \leq d$ is the Schmidt rank of the n 'th decomposition. Secondly, as discussed earlier, only a few terms are needed to accurately describe entanglement across bonds, thus allowing truncation of the S matrix (and likewise reduction of U and V) down to a chosen size D . Only the D largest singular values are kept, as these contribute the most to the state. Thus, the dimension of the matrices will cap at D : $(1 \times d), (d \times d^2), \dots, (d^n \times D), (D \times D), \dots, (D \times d^n) \dots (d \times 1)$, avoiding the otherwise exponential dimensional growth. [33]

5.4 Canonical Form

The MPS described in equation 89 is not unique, as writing

$$\tilde{A}^{j_n} = X_{n-1} A^{j_n} X_n^{-1} \quad (74)$$

describes the same state using different matrices. This gauge freedom allows for expressing the MPS in whichever way is most convenient often resulting in a much lower numerical cost of operations. By choosing a gauge the MPS is brought into a *canonical form*. [39]

Left-canonical matrix product state

The process above describing building the MPS from an arbitrary state produces an MPS in a *left-canonical* form. This means that all the matrices are left-normalized, such that

$$\sum_{j_n} A^{j_n \dagger} A^{j_n} = I. \quad (75)$$

This is a consequence of the matrix U (from the SVD) fulfilling $U^\dagger U = I$. Since the matrices A^{j_n} are reshaped from U , these properties persist.

$$\begin{aligned} \delta_{\alpha_n, \alpha'_n} &= \sum_{\alpha_{n-1} j_n} (U^\dagger)_{\alpha_n, (\alpha_{n-1} j_n)} U_{(\alpha_{n-1} j_n), \alpha'_n} \\ &= \sum_{\alpha_{n-1} j_n} (A^{j_n \dagger})_{\alpha_n, \alpha_{n-1}} A_{\alpha_{n-1}, \alpha'_n}^{j_n} \\ &= \sum_{j_n} \left(A^{j_n \dagger} A^{j_n} \right)_{\alpha_n, \alpha'_n} \end{aligned}$$

hence following the condition of equation 91. Figure 17 illustrates this, where two left-normalized matrices are contracted. This allows for contractions from the left without explicit calculation, which

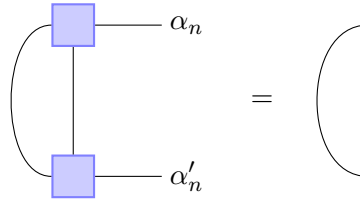


Figure 11: Contraction of two left-normalized A -matrices over their left index (shown as the arc) and their physical index (the connecting vertical line). The result is $\delta_{\alpha_n, \alpha'_n}$, adding the n 'th site to the contraction of all previous sites to the left.

is necessary when considering systems in the thermodynamic limit. In general $\sum_{j_n} A^{j_n} A^{j_n \dagger} \neq I$.

Right-canonical matrix product state

A right-canonical MPS can be built from 84 by starting from the other side of the chain. This implies multiplying U and S into the new Ψ -matrix, and reshaping $(V^\dagger)_{(\alpha_{n-1} j_n), \alpha_n}$ into matrices $B_{\alpha_{n-1}, \alpha_n}^{j_n}$. Hence, the right-canonical form of the MPS reads

$$|\psi\rangle = \sum_{j_1, \dots, j_N} B^{j_1} B^{j_2} \dots B^{j_{N-1}} B^{j_N} |j_1, \dots, j_N\rangle. \quad (76)$$

In this form all the matrices are right-normalized meaning

$$\sum_{j_n} B^{j_n} B^{j_n \dagger} = I, \quad (77)$$

which is illustrated in figure 18, which depict a contraction of right-normalized matrices at site n . Generally $\sum_{j_n} B^{j_n \dagger} B^{j_n} \neq I$.

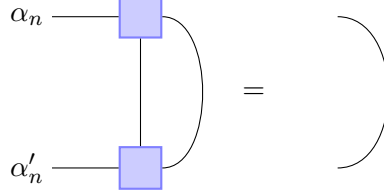


Figure 12: Contraction of two right-normalized B -matrices over their right index (shown as the arc) and their physical index (the connecting vertical line). The result is $\delta_{\alpha_n, \alpha'_n}$, adding the n 'th site to the contraction of all previous sites to the right.

Mixed-canonical matrix product state

The two above forms can also be mixed, which allows for easy bi-partitioning of the system into a Schmidt decomposition, which is useful for easy computations of reduced density matrices.

Consider the procedure of building an MPS in the left-canonical form as described in section 7.2, where the coefficients have been decomposed from the left up until site n

$$c_{j_1 \dots j_N} = \sum_{\alpha_n} (A^{j_1} \dots A^{j_n})_{\alpha_n} S_{\alpha_n, \alpha_n} (V^\dagger)_{\alpha_n, (j_{n+1} \dots j_N)}. \quad (78)$$

From here the V^\dagger can be reshaped into the matrix $\Psi_{(\alpha_n j_{n+1} \dots j_{N-1}), j_N}$, which can then be successively decomposed from the right, resulting in a set of right-normalized matrices. The final result reads

$$c_{j_1 \dots j_N} = A^{j_1} \dots A^{j_n} S B^{j_{n+1}} B^{j_N}, \quad (79)$$

which is illustrated in figure 19. In this form the Schmidt decomposition can be read directly of the

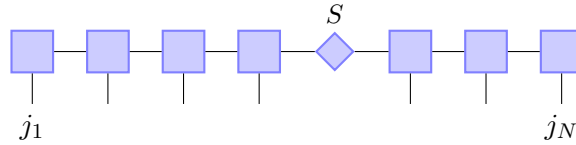


Figure 13: Diagrammatic representation of an MPS in mixed-canonical form, where the tensors on the left are left-normalized, while the tensors on the right are right-normalised. The matrix S is a diagonal, singular valued matrix.

MPS by introducing the vectors

$$|\alpha_n\rangle_A = \sum_{j_1, \dots, j_n} (A^{j_1} \dots A^{j_n})_{1, \alpha_n} |j_1, \dots, j_n\rangle \quad (80)$$

$$|\alpha_n\rangle_B = \sum_{j_{n+1}, \dots, j_N} (B^{j_{n+1}} \dots B^{j_N})_{\alpha_n, 1} |j_{n+1}, \dots, j_N\rangle \quad (81)$$

whereby the state can be written in the form

$$|\psi\rangle = \sum_{\alpha_n} S_{\alpha_n, \alpha_n} |\alpha_n\rangle_A |\alpha_n\rangle_B. \quad (82)$$

In order for this to be a Schmidt decomposition $\sum_{\alpha_n} (S_{\alpha_n, \alpha_n})^2 = 1$, which is fulfilled from the properties of the SVD, and the states on A and B have to be orthonormal respectively, which they are by construction.

Bringing a matrix product state into canonical form

So far it has only been presented how to bring an MPS into a canonical form when building it. However, an arbitrary, pre-built MPS can be brought into a canonical form through a series of SVD's, again exploiting the unitarity or left-/right-normalization of the resulting matrices.

Consider a general MPS

$$|\psi\rangle = \sum_{j_1, \dots, j_N} \sum_{\alpha_1, \dots} M_{1, \alpha_1}^{j_1} M_{\alpha_1, \alpha_2}^{j_2} M_{\alpha_2, \alpha_3}^{j_3} \dots |j_1, \dots, j_N\rangle, \quad (83)$$

which has to be brought into a left-canonical form. By grouping the physical and left (row) index of $M_{1, \alpha_1}^{j_1}$, one can bring this set of matrices into the form of a single matrix, where an SVD can be applied to it. This yields $M = ASV^\dagger$, where $A^\dagger A = I$, such that A is left-normalized as desired.

$$\begin{aligned} & \sum_{j_1, \dots, j_N} \sum_{\alpha_1, \dots} M_{(j_1, 1), \alpha_1} M_{\alpha_1, \alpha_2}^{j_2} M_{\alpha_2, \alpha_3}^{j_3} \dots |j_1, \dots, j_N\rangle \\ &= \sum_{j_1, \dots, j_N} \sum_{\alpha_1, \dots} \sum_{s_1} A_{(j_1, 1), s_1} S_{s_1, s_1} V_{s_1, \alpha_1}^\dagger M_{\alpha_1, \alpha_2}^{j_2} M_{\alpha_2, \alpha_3}^{j_3} \dots |j_1, \dots, j_N\rangle \\ &= \sum_{j_1, \dots, j_N} \sum_{\alpha_1, \dots} \sum_{s_1} A_{1, s_1}^{j_1} \left(S_{s_1, s_1} V_{s_1, \alpha_1}^\dagger M_{\alpha_1, \alpha_2}^{j_2} \right) M_{\alpha_2, \alpha_3}^{j_3} \dots |j_1, \dots, j_N\rangle \\ &= \sum_{j_1, \dots, j_N} \sum_{\alpha_2, \dots} \sum_{s_1} A_{1, s_1}^{j_1} \tilde{M}_{s_1, \alpha_2}^{j_2} M_{\alpha_2, \alpha_3}^{j_3} \dots |j_1, \dots, j_N\rangle, \end{aligned} \quad (84)$$

where $\tilde{M}_{s_1, \alpha_2}^{j_2} = \sum_{\alpha_1} S_{s_1, s_1} V_{s_1, \alpha_1}^\dagger M_{\alpha_1, \alpha_2}^{j_2}$. This procedure can be iterated through the entire chain leaving an MPS in the left-canonical form.

Likewise, an MPS can also be brought into a right-canonical form by grouping the physical index with the right (column) index of M yielding the SVD $M = USB$, where $BB^\dagger = I$. Iterating through the chain from the right will thus make the MPS right-canonical. [?]

5.5 Overlaps and Efficient Contractions

Once the MPS is built it is important to understand how to use it for efficient calculations of overlaps and expectation values. For this, consider the general case of two states $|\psi\rangle$ and $|\phi\rangle$ described by the matrices M and \tilde{M} . The overlap between these states reads

$$\langle\phi|\psi\rangle = \sum_{j_1, \dots, j_N} \tilde{M}^{j_1*} \dots \tilde{M}^{j_N*} M^{j_1} \dots M^{j_N} = \sum_{j_1, \dots, j_N} \tilde{M}^{j_N\dagger} \dots \tilde{M}^{j_1\dagger} M^{j_1} \dots M^{j_N}, \quad (85)$$

which is represented grafically in figure 20. The evaluation of the overlap can be drastically sped up by

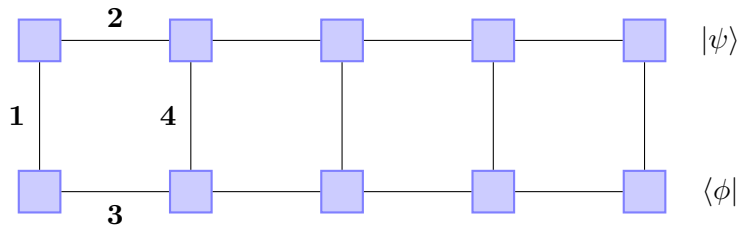


Figure 14: Efficient contraction of the overlap between two general states $|\psi\rangle$ and $|\phi\rangle$. By contracting the bonds in the specified order the computational cost is greatly reduced.

considering the optimal order of contractions, which corresponds to an optimal bracketing of equation 101:

$$\langle\phi|\psi\rangle = \sum_{j_N} \tilde{M}^{j_N\dagger} \left(\dots \left(\sum_{j_2} \tilde{M}^{j_2\dagger} \left(\sum_{j_1} \tilde{M}^{j_1\dagger} M^{j_1} \right) M^{j_2} \right) \dots \right) M^{j_N}. \quad (86)$$

The innermost bracket a matrix is formed from the multiplication of a column and a row vector and the first physical index, j_1 , is summed over. In the second set of brackets three matrices are multiplied and the second physical index is summed over. This procedure continues, however, after the first contraction the complexity of the operation does not increase. This is in contrast to the scenario, where an arbitrary bond was contracted over at first. If d is the dimension of the Hilbert spaces at the physical sites, and all the matrices are of dimension $(D \times D)$, then the total operational cost of the optimal contraction is $O(ND^3d)$ [?]. The order of contractions is illustrated in figure 101.

In the special case of calculating a norm, $\langle \psi | \psi \rangle$ having brought the MPS to a canonical form greatly simplifies the calculation, as having an either left- or right-canonical MPS implies a norm of 1. This can easily be seen for left-canonical form when considering equation 91 and 102. Due to the left-orthogonality every matrix product is I , causing it to drop out. Finally, summing over the final index yields 1.

5.6 Matrix Product Operators (MPO)

An MPO is an operator expressed in the formalism of an MPS. Thus, operators can easily be incorporated as part of the tensor networks, where they are evaluated by contraction over their bonds. Consider a single coefficient of an MPS

$$\langle \mathbf{j} | \psi \rangle = M^{j_1} M^{j_2} \dots M^{j_N} . \quad (87)$$

Expressing an operator \hat{O} in the basis of the local states, one can write it in a similar manner

$$\hat{O} = \sum_{\mathbf{j}, \mathbf{j}'} |\mathbf{j}\rangle \langle \mathbf{j} | \hat{O} | \mathbf{j}' \rangle \langle \mathbf{j}'| = \sum_{\mathbf{j}, \mathbf{j}'} W^{j_1, j'_1} W^{j_2, j'_2} \dots W^{j_N, j'_N} |\mathbf{j}\rangle \langle \mathbf{j}'| , \quad (88)$$

where the coefficients are $\langle \mathbf{j} | \hat{O} | \mathbf{j}' \rangle = W^{j_1, j'_1} W^{j_2, j'_2} \dots W^{j_N, j'_N}$. The matrices W^{j_n, j'_n} are just like the M -matrices, except the the representation of the operators needs both an ingoing and an outgoing physical index. This corresponds graphically to having two vertical lines; one corresponding to the ingoing physical state, the other corresponding to the outgoing physical state. A pictorial representation of the operator \hat{O} can be seen in figure 21.

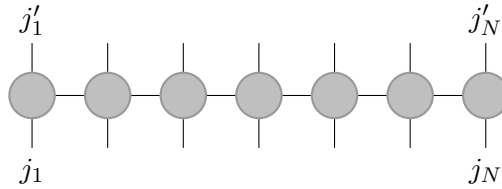


Figure 15: An operator \hat{O} expressed in the MPS form (MPO). The resulting matrix product has two vertical lines corresponding to an ingoing and outgoing physical state.

Applying an MPO to an MPS

Applying a matrix product operator to a matrix product state is simply a matrix multiplication, where the matching physical indices are summed over:

$$\begin{aligned}
 \hat{O} |\psi\rangle &= \sum_{\mathbf{j}, \mathbf{j}'} \left(M^{j'_1} M^{j'_2} \dots \right) \left(W^{j'_1, j_1} W^{j'_2, j_2} \dots \right) |\mathbf{j}\rangle \\
 &= \sum_{\mathbf{j}, \mathbf{j}'} \sum_{\alpha, \beta} \left(M^{j'_1}_{1, \alpha_1} M^{j'_2}_{\alpha_1, \alpha_2} \dots \right) \left(W^{j'_1, j_1}_{1, \beta_1} W^{j'_2, j_2}_{\beta_1, \beta_2} \dots \right) |\mathbf{j}\rangle \\
 &= \sum_{\mathbf{j}, \mathbf{j}'} \sum_{\alpha, \beta} \left(M^{j'_1}_{1, \alpha_1} W^{j'_1, j_1}_{1, \beta_1} \right) \left(M^{j'_2}_{\alpha_1, \alpha_2} W^{j'_2, j_2}_{\beta_1, \beta_2} \right) \dots |\mathbf{j}\rangle \\
 &= \sum_{\mathbf{j}} \sum_{\alpha, \beta} N^{j_1}_{(1,1), (\alpha_1, \beta_1)} N^{j_2}_{(\alpha_1, \beta_1), (\alpha_2, \beta_2)} \dots |\mathbf{j}\rangle \\
 &= \sum_{\mathbf{j}} N^{j_1} N^{j_2} \dots |\mathbf{j}\rangle = |\phi\rangle
 \end{aligned} \tag{89}$$

The result is a new MPS, $|\phi\rangle$, which is described by the matrices N^{j_n} . These matrices have the dimensions of the product of the dimensions of the original MPS and MPO. Thus, applying an operator leaves the form of the MPS invariant at a cost of increased matrix dimensions, which is illustrated in figure 22. If the dimension of the MPS is D , while the dimension of the MPO is D_W , the total

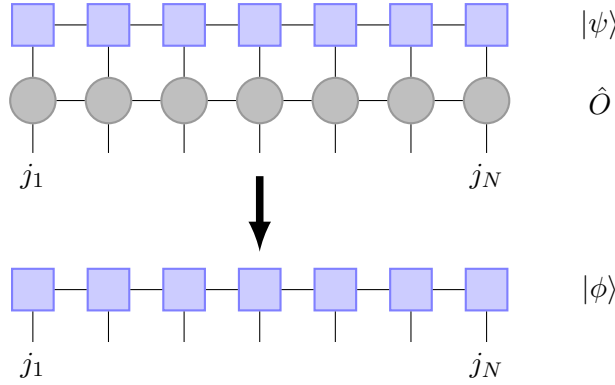


Figure 16: Application of an MPO, \hat{O} , on an MPS, $|\psi\rangle$. Matching physical indices are contracted resulting in a new MPS, $|\phi\rangle$, with increased matrix dimensions.

computational cost of the operation is $O(Nd^2D_W^2D^2)$. [?, 25]

Efficient evaluation of local operators

Often one only needs to apply an operator to a single site. Consider the local operator $\hat{O}^{[n]}$, where the square brackets denote the origin of the operator. As $\hat{O}^{[n]}$ is local it only acts on site n and can be expressed in the basis of said site

$$\hat{O}^{[n]} = \sum_{j_n, j'_n} O^{j_n, j'_n} |j_n\rangle \langle j'_n|. \tag{90}$$

Applying this operator to a state describing the entire system implies applying the identity operator on all other sites such that

$$\hat{O} = I^{[1]} \otimes I^{[2]} \otimes \dots \otimes O^{[n]} \otimes \dots \otimes I^{[N]}. \tag{91}$$

In order to evaluate the expectation value of an operator using an MPS the operator is expressed as a product of matrices $O^{j_1, j'_1} \dots O^{j_N, j'_N}$. In the case of a single local operator most of these matrices

will be identities. This matrix product can then be multiplied with the matrices of the MPS. Again applying optimal bracketing simplifies the computation:

$$\langle \psi | \hat{O} | \psi \rangle = \sum_{j_1, \dots, j_N} \sum_{j'_1, \dots, j'_N} \tilde{M}^{j_1*} \dots \tilde{M}^{j_N*} O^{j_1, j'_1} \dots O^{j_N, j'_N} M^{j'_1} \dots M^{j'_N} \quad (92)$$

$$= \sum_{j_N, j'_N} O^{j_N, j'_N} \tilde{M}^{j_N \dagger} \left(\dots \left(\sum_{j_2, j'_2} O^{j_2, j'_2} \tilde{M}^{j_2 \dagger} \left(\sum_{j_1, j'_1} O^{j_1, j'_1} \tilde{M}^{j_1 \dagger} M^{j'_1} \right) M^{j'_2} \right) \dots \right) M^{j'_N} . \quad (93)$$

Although this operation sums over two indices in every brackets most of the matrices O^{j_n, j'_n} are identities causing them to drop out, thus reducing the sum to a single index for most sites. Hence the operational cost is essentially still $O(ND^3d)$. [?]

However, the above calculation can be simplified even further when considering canonical forms. Consider an MPS in the mixed-canonical form, where all matrices left of site n are left-normalized, while all matrices to the right are right-normalized. In this case the two sides can be contract without any calculation, resulting in the network shown in figure 23, which reduces the calculation to simply

$$\langle \psi | \hat{O}^{[n]} | \psi \rangle = \sum_{j_n, j'_n} O^{j_n, j'_n} \text{Tr} \left(M^{j_n \dagger} M^{j'_n} \right) \quad (94)$$

with the computational cost $O(D^2d^2)$.

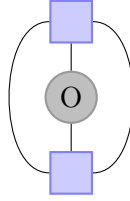


Figure 17: Contracted network for evaluating $\langle \psi | \hat{O}^{[n]} | \psi \rangle$, where matrices are left- and right-normalized to the left and right of site n .

5.7 Correlation Functions

Correlation length

6 Calculating Ground States with MPS

For large systems exact diagonalization of the Hamiltonian is impossible, whereby one must resort to variational methods in order to find ground states. This involves finding the MPS $|\psi\rangle$, which minimizes

$$E = \frac{\langle \psi | \hat{H} | \psi \rangle}{\langle \psi | \psi \rangle} . \quad (95)$$

The most efficient way to do this, is to express the Hamiltonian as an MPO as described in equation 104, which makes it easy to apply to an MPS. While this may seem difficult at first, it can actually be done rather easily without any lengthy calculations. To see this consider the following example:

Example: Building an MPO from a Hamiltonian

Consider a Hamiltonian consisting of next-neighbour interaction terms and a field term

$$\hat{H} = \sum_i \frac{J}{2} \hat{S}_i^+ \hat{S}_{i+1}^- + \frac{J}{2} \hat{S}_i^- \hat{S}_{i+1}^+ + J^z \hat{S}_i^z \hat{S}_{i+1}^z - h \hat{S}_i^z , \quad (96)$$

which can be expressed as a sum of strings of operators - most of these being identities just like in equation 106. Moving through such a string from the right one will at some point encounter one of 4 non-trivial operators. This can be summarized as 5 different possible states of the string of operators on a given bond:

1. Only identities to the right of the bond.
2. An \hat{S}^+ operator just to the right of the bond.
3. An \hat{S}^- operator just to the right of the bond.
4. An \hat{S}^z operator just to the right of the bond.
5. A completed interaction *or* the field term, $h\hat{S}^z$, somewhere to the right.

When moving through the string of operators from the right, only certain transitions between these state are possible. For instance $\mathbf{1} \rightarrow \mathbf{1}$, where one starts in state 1, and the next operator is an identity. Likewise, $\mathbf{1} \rightarrow \mathbf{2}, \mathbf{3}, \mathbf{4}, \mathbf{5}$ are all possible, since these transitions represent the next operator being one of the interactions or the field. Next, $\mathbf{2} \rightarrow \mathbf{5}$ completes the interaction $\frac{J}{2}\hat{S}^+\hat{S}^-$ - similar transitions exists for all the possible interactions $\mathbf{3}, \mathbf{4} \rightarrow \mathbf{5}$. Finally, the transition $\mathbf{5} \rightarrow \mathbf{5}$ is needed to continue iterating through the operator chain after having passed the interactions. This can be encoded in the operator-valued matrix

$$W^{[i]} = \begin{pmatrix} \hat{I} & 0 & 0 & 0 & 0 \\ \hat{S}^+ & 0 & 0 & 0 & 0 \\ \hat{S}^- & 0 & 0 & 0 & 0 \\ \hat{S}^z & 0 & 0 & 0 & 0 \\ -h\hat{S}^z & (J/2)\hat{S}^- & (J/2)\hat{S}^+ & J^z\hat{S}^z & \hat{I} \end{pmatrix}, \quad (97)$$

which contains all the allowed transitions between the five state [?]. When one starts moving through the operator chain from the right side, one obviously begins in state 1 and ends in state 5. This can be encoded in the two vectors

$$\vec{v}_{left} = (0, 0, 0, 0, 1) \quad , \quad \vec{v}_{right} = (1, 0, 0, 0, 0)^T.$$

Thus, the Hamiltonian described in equation 112 can be written as an MPO

$$\hat{H} = \sum_{\substack{j_1, \dots, j_N \\ j'_1, \dots, j'_N}} \vec{v}_{left} W^{[1]j_1, j'_1} W^{[2]j_2, j'_2} \dots W^{[N]j_N, j'_N} \vec{v}_{right} |j_1, \dots, j_N\rangle \langle j'_1, \dots, j'_N|. \quad (98)$$

Observe how all of this was done without having to do a single numerical computation. The resulting MPO can now readily be applied to an MPS.

This is just an example for a simple Hamiltonian with only next-neighbour interactions. Interactions of longer range require intermediate states consisting of identities in order to connect the interacting operators. However, doing this is fairly simple and will often result in MPO's of a similar low dimension as the one shown in the example.

6.1 Efficient Application of a Hamiltonian to an MPS

In section 7.5 it was explained how to apply an MPO to an MPS, and it was shown how to efficiently evaluate local operators. While Hamiltonians may contains local terms, they are rarely completely local, and since $\hat{H}|\psi\rangle$ must be evaluated many times during variational methods, the naive approach

of section 7.5 simply doesn't cut it.

Consider an MPS in a mixed-canonical form

$$\begin{aligned} |\psi\rangle &= \sum_{\mathbf{j}} A^{j_1} \dots A^{j_{n-1}} \Psi^{j_n} B^{j_{n+1}} \dots B^{j_N} \\ &= \sum_{\alpha_{n-1}, \alpha_n} |\alpha_{n-1}\rangle_A \Psi_{\alpha_{n-1}, \alpha_n}^{j_n} |\alpha_n\rangle_B, \end{aligned} \quad (99)$$

where the set of matrices at the n 'th site has not been brought into either canonical form, and a Hamiltonian in the same form as described in equation 114. For notations sake the opening and closing vectors have been combined with the outer matrices of the MPO. For the variational search of the ground state huge reductions in computational cost can be found by re-using as many calculations as possible. Consider in the basis $\{|\alpha_{n-1}\rangle, |j_n\rangle, |\alpha_n\rangle\}$ the individual matrix elements of the Hamiltonian

$$\langle \alpha_{n-1} j_n \alpha_n | \hat{H} | \alpha'_{n-1} j'_n \alpha'_n \rangle = \sum_{\mathbf{j}, \mathbf{j}'} W^{j_1, j'_1} \dots W^{j_N, j'_N} \langle \alpha_{n-1} j_n \alpha_n | \mathbf{j} \rangle \langle \mathbf{j}' | \alpha'_{n-1} j'_n \alpha'_n \rangle$$

Since the basis of local states $\{|\mathbf{j}\rangle\}$ shares a state with the basis $\{|\alpha_{n-1}\rangle, |j_n\rangle, |\alpha_n\rangle\}$, the above expression can be re-written using $\sum_{\mathbf{j}} \langle j_n | \mathbf{j} \rangle = \sum_{\mathbf{j}^*} |\mathbf{j}^*\rangle$, where " $*$ " means "excluding j_n ". Thus,

$$\begin{aligned} &\langle \alpha_{n-1} j_n \alpha_n | \hat{H} | \alpha'_{n-1} j'_n \alpha'_n \rangle \\ &= \sum_{\mathbf{j}^*, \mathbf{j}'^*} W^{j_1, j'_1} \dots W^{j_n, j'_n} \dots W^{j_N, j'_N} \\ &\quad \times \langle \alpha_{n-1} | j_1 \dots j_{n-1} \rangle \langle \alpha_n | j_{n+1} \dots j_N \rangle \langle j'_1 \dots j'_{n-1} | \alpha'_{n-1} \rangle \langle j'_{n+1} \dots j'_N | \alpha'_n \rangle \\ &= \sum_{\mathbf{j}^*, \mathbf{j}'^*} W^{j_1, j'_1} \dots W^{j_n, j'_n} \dots W^{j_N, j'_N} \\ &\quad \times (A^{j_1} \dots A^{j_{n-1}})_{1, \alpha_{n-1}}^* (B^{j_{n+1}} \dots B^{j_N})_{\alpha_n, 1}^* (A^{j'_1} \dots A^{j'_{n-1}})_{1, \alpha'_{n-1}} (B^{j'_{n+1}} \dots B^{j'_N})_{\alpha'_n, 1} \\ &= \sum_{\alpha_n, \beta_n, \alpha'_n} \left(\sum_{j_1, j'_1} A_{1, \alpha_1}^{j_1*} W_{1, \beta_1}^{j_1, j'_1} A_{1, \alpha'_1}^{j'_1} \right) \left(\sum_{j_2, j'_2} A_{\alpha_1, \alpha_2}^{j_2*} W_{\beta_1, \beta_2}^{j_2, j'_2} A_{\alpha'_1, \alpha'_2}^{j'_2} \right) \dots W_{\beta_{n-1}, \beta_n}^{j_n, j'_n} \\ &\quad \times \left(\sum_{j_{n+1}, j'_{n+1}} B_{\alpha_n, \alpha_{n+1}}^{j_{n+1}*} W_{\beta_n, \beta_{n+1}}^{j_{n+1}, j'_{n+1}} B_{\alpha'_n, \alpha'_{n+1}}^{j'_{n+1}} \right) \left(\sum_{j_N, j'_N} B_{\alpha_{N-1}, 1}^{j_N*} W_{\beta_{N-1}, 1}^{j_N, j'_N} B_{\alpha'_{N-1}, 1}^{j'_N} \right). \end{aligned}$$

While this expression may seem terrible complicated due to all the indices, it is actually rather easy to understand. First, the matrix element is written excluding the local basis state $|j_n\rangle$. Next, the Hamilton MPO is projected into the block states of A and B, $|\alpha_{n-1}\rangle_A$ and $|\alpha_n\rangle_B$, which can easily be seen when considering equations 96 and 97. Finally, the matrices are grouped according to their expansion in the local basis. Working with the above expression appears cumbersome, but as can be seen in figure 24, what has been done is simply a decoupling of the system into three distinct parts. Since both the left and right side of the network is connected, one can contract these parts into two separate tensors L and R called "environments":

$$L_{\alpha_{n-1}, \beta_{n-1}, \alpha'_{n-1}} = \sum_{\substack{\{\alpha_i, \beta_i, \alpha'_i\} \\ i < n-1}} \left(\sum_{j_1, j'_1} A_{1, \alpha_1}^{j_1*} W_{1, \beta_1}^{j_1, j'_1} A_{1, \alpha'_1}^{j'_1} \right) \dots \left(\sum_{j_{n-1}, j'_{n-1}} A_{\alpha_{n-2}, \alpha_{n-1}}^{j_{n-1}*} W_{\beta_{n-2}, \beta_{n-1}}^{j_{n-1}, j'_{n-1}} A_{\alpha'_{n-2}, \alpha'_{n-1}}^{j'_{n-1}} \right) \quad (100)$$

$$R_{\alpha_n, \beta_n, \alpha'_n} = \sum_{\substack{\{\alpha_i, \beta_i, \alpha'_i\} \\ i > n}} \left(\sum_{j_{n+1}, j'_{n+1}} B_{\alpha_n, \alpha_{n+1}}^{j_{n+1}*} W_{\beta_n, \beta_{n+1}}^{j_{n+1}, j'_{n+1}} B_{\alpha'_n, \alpha'_{n+1}}^{j'_{n+1}} \right) \left(\sum_{j_N, j'_N} B_{\alpha_{N-1}, 1}^{j_N*} W_{\beta_{N-1}, 1}^{j_N, j'_N} B_{\alpha'_{N-1}, 1}^{j'_N} \right) \quad (101)$$

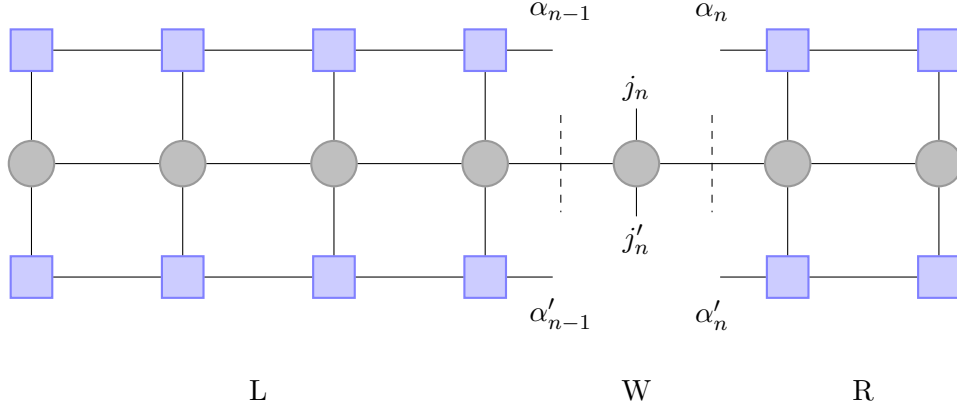


Figure 18: Representation of the matrix element $\langle \alpha_{n-1} j_n \alpha_n | \hat{H} | \alpha'_{n-1} j'_n \alpha'_n \rangle$ as an MPS/MPO network. Expressing the matrix element in this form decouples the network in three parts: The matrices of the MPO, $W^{[n]}$, connecting the physical sites of the matrix element, and contracted parts L and R consisting of the rest of the MPO and the part of the MPS in left- and right-canonical form respectively.

From these contractions the obvious tripartite structure of the Hamiltonian matrix element, which could be seen in figure 24, can be written in a compact way

$$\langle \alpha_{n-1} j_n \alpha_n | \hat{H} | \alpha'_{n-1} j'_n \alpha'_n \rangle = \sum_{\beta_{n-1}, \beta_n} L_{\alpha_{n-1}, \beta_{n-1}, \alpha'_{n-1}} W_{\beta_{n-1}, \beta_n}^{j_n, j'_n} R_{\alpha_n, \beta_n, \alpha'_n} . \quad (102)$$

Finally, applying this parametrization of the Hamiltonian to the MPS described in equation 115 yields [?]

$$\hat{H} |\psi\rangle = \sum_{\beta_{n-1}, \beta_n} \sum_{\alpha'_{n-1}, j'_n, \alpha'_n} L_{\alpha_{n-1}, \beta_{n-1}, \alpha'_{n-1}} W_{\beta_{n-1}, \beta_n}^{j_n, j'_n} R_{\alpha_n, \beta_n, \alpha'_n} \Psi_{\alpha'_{n-1}, \alpha'_n}^{j'_n} |\alpha_{n-1}\rangle_A |j_n\rangle |\alpha_n\rangle_B . \quad (103)$$

As mentioned earlier, evaluating $\hat{H} |\psi\rangle$ must be done many times during a variational search of the ground state, hence this operation must be executed as fast as possible. Examining equation 119 one will notice, that while the boundaries of L and R will change depending on which site is being optimized, the bulk of the two tensors remain constant through a lot of the calculations. Instead of calculating L and R from equations 116 and 117 explicitly every time equation 119 has to be evaluated, one can iteratively build them, since they only change by one column of the network at a time. This allows reuse of a lot of information resulting in much lower computational cost.

Consider the construction of the tensor $L^{[i]}$. This can be built iteratively from the left by contracting the previous left-tensor $L^{[i-1]}$ with the i 'th column of the network consisting of $A^{[i]}$, $W^{[i]}$ and $A^{[i]\dagger}$

$$L_{\alpha_i, \beta_i, \alpha'_i}^{[i]} = \sum_{\substack{j_i, j'_i \\ \alpha_{i-1}, \beta_{i-1}, \alpha'_{i-1}}} W_{\beta_{i-1}, \beta_i}^{[i] j_i, j'_i} \left(A^{[i] j_i \dagger} \right)_{\alpha_i, \alpha_{i-1}} L_{\alpha_{i-1}, \beta_{i-1}, \alpha'_{i-1}}^{[i-1]} A_{\alpha'_{i-1}, \alpha'_i}^{[i] j'_i} . \quad (104)$$

This iterative update of $L^{[i]}$ can be seen illustrated in figure 25. The square-bracket notation has been re-introduced to keep track of the tensors relation to the physical sites. In order to remain consistent with notation, the dummy scalars $L_{\alpha_0, \beta_0, \alpha'_0}^{[0]} = 1 = \alpha_0, \beta_0, \alpha'_0$ has been introduced. Performing this update can be sped up when considering the properties of the Hamiltonian and the canonical form. Looking at matrix representation, $W^{[i]}$, of the example Hamiltonian in equation 113 one will notice that most entries are zeroes. While this may not be the case for every Hamiltonian, knowing the structure of the corresponding MPO allows one to simplify multiplications. Furthermore, for the last index in the $W^{[i]}$ matrix, $\beta_i = D_W$, only identities will operate towards the left, if the MPO is constructed according to the rules outlined earlier. Since the matrices $A^{[i]}$ are left normalized, this implies that $L_{\alpha_i, D_W, \alpha'_i}^{[i]} = \delta_{\alpha_i, \alpha'_i}$, greatly simplifying calculations.

It is important to store every iteration of $L^{[i]}$, since L will grow and shrink constantly throughout the

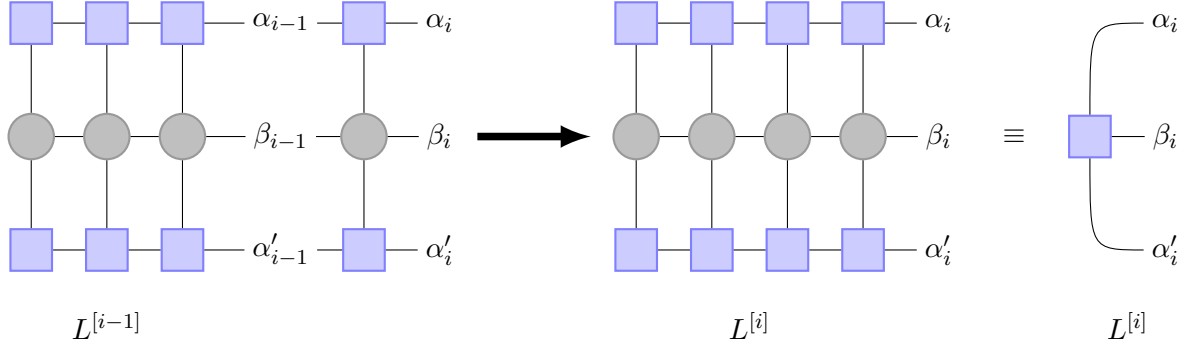


Figure 19: Iterative update from $L^{[i-1]}$ to $L^{[i]}$. This is done through a contraction of $L^{[i-1]}$ with $A^{[i]}$, $W^{[i]}$ and $A^{[i]†}$. The result is a tensor with three horizontal legs.

variational search of the ground state, whereby every iteration of $L^{[i]}$ will be used multiple times. By keeping track of every iteration, an update of the MPS at the j 'th site will only affect $L^{[i]}$ for $i \geq j$ - and this update can be done swiftly using the previously stored iterations.

Naturally all of this applies when building the right tensor R . Here one starts from the right and moves left when iteratively contracting the tensor. Likewise, right-normalization simplify the contractions when $b_i = 1$, implying only identities to the left in the chain of operators.

Finally, a re-bracketing for optimal matrix multiplication allows for an even lower operational cost

$$L_{\alpha_i, \beta_i, \alpha'_i}^{[i]} = \sum_{j_i, \alpha_{i-1}} \left(A^{[i]j_i \dagger} \right)_{\alpha_i, \alpha_{i-1}} \left(\sum_{j'_i, \beta_{i-1}} W_{\beta_{i-1}, \beta_i}^{[i]j_i, j'_i} \left(\sum_{\alpha'_{i-1}} L_{\alpha_{i-1}, \beta_{i-1}, \alpha'_{i-1}}^{[i-1]} A_{\alpha'_{i-1}, \alpha'_i}^{[i]j'_i} \right) \right)$$

$$\hat{H} |\psi\rangle = \sum_{\beta_{n-1}, \alpha'_{n-1}} L_{\alpha_{n-1}, \beta_{n-1}, \alpha'_{n-1}} \left(\sum_{\beta_n, j'_n} W_{\beta_{n-1}, \beta_n}^{j_n, j'_n} \left(\sum_{\alpha'_n} R_{\alpha_n, \beta_n, \alpha'_n} \Psi_{\alpha'_{n-1}, \alpha'_n}^{j'_n} \right) \right) |\alpha_{n-1}\rangle_A |j_n\rangle |\alpha_n\rangle_B ,$$

where they both scale as $O(dD^3D_W)$.

6.2 Iterative Ground State Search

In order to find the ground state of the system one can introduce a Lagrangian multiplier and extremize

$$\langle \psi | \hat{H} | \psi \rangle - \lambda \langle \psi | \psi \rangle , \quad (105)$$

whereby the desired ground state, $|\psi\rangle$, and ground state energy, λ^0 , will be reached.

Consider an MPS of the form $M^{[1]} \dots M^{[N]}$. Trying to optimize the entire MPS at the same time is a highly non-linear problem involving an extremely large number of variables. However, the problem can be linearised by only considering the matrix entries $M_{\alpha_{n-1}, \alpha_n}^{j_n}$ at site n as variables, while keeping the rest of the MPS constant. By varying the matrix elements one site at a time, one will continuously find states lower in energy until convergence is reached. However, this procedure is very prone to getting stuck in a local extrema. To circumvent this one can consider two sites at a time and optimize with regards to a tensor, created by momentarily merging the two sites.^[41]

Considering equation 121 one has to calculate

$$\langle \psi | \hat{H} | \psi \rangle = \sum_{\substack{j_n, j'_n \\ j_{n+1}, j'_{n+1}}} \sum_{\substack{\alpha'_{n-1}, \alpha'_n, \alpha'_{n+1} \\ \alpha_{n-1}, \alpha_n, \alpha_{n+1}}} \sum_{\beta_{n-1}, \beta_n, \beta_{n+1}} L_{\alpha_{n-1}, \beta_{n-1}, \alpha'_{n-1}}^{[n-1]} W_{\beta_{n-1}, \beta_n}^{[n]j_n, j'_n} W_{\beta_n, \beta_{n+1}}^{[n+1]j_{n+1}, j'_{n+1}} \\ \times R_{\alpha_{n+1}, \beta_{n+1}, \alpha'_{n+1}}^{[n+2]} M_{\alpha_{n-1}, \alpha_n}^{[n]j_n} M_{\alpha'_{n-1}, \alpha'_n}^{[n]j'_n*} M_{\alpha_n, \alpha_{n+1}}^{[n+1]j_{n+1}} M_{\alpha'_n, \alpha'_{n+1}}^{[n+1]j'_{n+1}*} \quad (106)$$

$$\langle \psi | \psi \rangle = \sum_{j_n, j_{n+1}} \sum_{\substack{\alpha'_{n-1} \\ \alpha'_n, \alpha'_{n+1}}} \sum_{\substack{\alpha_{n-1}, \alpha_n \\ \alpha_{n+1}}} \Psi_{\alpha_{n-1}, \alpha'_{n-1}}^A M_{\alpha_{n-1}, \alpha_n}^{[n]j_n} M_{\alpha'_{n-1}, \alpha'_n}^{[n]j'_n*} M_{\alpha_n, \alpha_{n+1}}^{[n+1]j_{n+1}} M_{\alpha'_n, \alpha'_{n+1}}^{[n+1]j'_{n+1}*} \Psi_{\alpha_{n+1}, \alpha'_{n+1}}^B, \quad (107)$$

where the Hamiltonian from equation 119 has been re-ordered to accommodate for examining two sites, n and $n+1$, at a time, and

$$\Psi_{\alpha_{n-1}, \alpha'_{n-1}}^A = \sum_{j_1, \dots, j_{n-1}} \left(M^{j_{n-1}\dagger} \dots M^{j_1\dagger} M^{j_1} \dots M^{j_{n-1}} \right)_{\alpha_{n-1}, \alpha'_{n-1}} \quad (108)$$

$$\Psi_{\alpha_{n+1}, \alpha'_{n+1}}^B = \sum_{j_{n+2}, \dots, j_N} \left(M^{j_{n+2}} \dots M^{j_N} M^{j_N\dagger} \dots M^{j_{n+2}\dagger} \right)_{\alpha'_{n+1}, \alpha_{n+1}}. \quad (109)$$

Further simplifications can be made for mixed-canonical forms, if sites 1 through $n-1$ are left-normalized, and sites $n+2$ through N are right-normalized

$$\Psi_{\alpha_{n-1}, \alpha'_{n-1}}^A = \delta_{\alpha_{n-1}, \alpha'_{n-1}} \quad , \quad \Psi_{\alpha_{n+1}, \alpha'_{n+1}}^B = \delta_{\alpha_{n+1}, \alpha'_{n+1}}. \quad (110)$$

From here one has to find the extremum of equation 121 with respect to $M_{\alpha'_{n-1}, \alpha'_n}^{[n]j'_n*} M_{\alpha'_n, \alpha'_{n+1}}^{[n+1]j'_{n+1}*}$. Looking at equation 122 and 123 this may seem like a daunting task, however, it can actually be done fairly easily through the following sequence:

Two-site update for iterative ground state search

1. **Merge:** Contract the two matrices $M^{[n]}$ and $M^{[n+1]}$ over the bond α_n creating a two-site tensor

$$\Theta_{\alpha_{n-1}, \alpha_{n+1}}^{j_n, j_{n+1}} = \sum_{\alpha_n} M_{\alpha_{n-1}, \alpha_n}^{[n]j_n} M_{\alpha_n, \alpha_{n+1}}^{[n+1]j_{n+1}} \quad (111)$$

2. **Solve eigenproblem:** The problem is simply an eigenvalue problem, which can be seen by reshaping

$$H_{(\alpha_{n-1}j_nj_{n+1}, \alpha_{n+1}), (\alpha'_{n-1}j'_nj'_{n+1}, \alpha'_{n+1})} = \\ = \sum_{\substack{\beta_{n-1}, \beta_n \\ \beta_{n+1}}} L_{\alpha_{n-1}, \beta_{n-1}, \alpha'_{n-1}}^{[n-1]} W_{\beta_{n-1}, \beta_n}^{[n]j_n, j'_n} W_{\beta_n, \beta_{n+1}}^{[n+1]j_{n+1}, j'_{n+1}} R_{\alpha_{n+1}, \beta_{n+1}, \alpha'_{n+1}}^{[n+2]} \quad (112)$$

$$v_{\alpha_{n-1}j_nj_{n+1}\alpha_{n+1}} = \Theta_{\alpha_{n-1}, \alpha_{n+1}}^{j_n, j_{n+1}} \quad (113)$$

such that

$$Hv - \lambda v = 0. \quad (114)$$

Solving this for the lowest eigenvalue λ_0 yields $v_{\alpha_{n-1}j_nj_{n+1}\alpha_{n+1}}^0$, which can be reshaped back to the now optimized $\tilde{\Theta}_{\alpha_{n-1}, \alpha_{n+1}}^{j_n, j_{n+1}}$.

3. **Unmerge:** Reshape the updated $\tilde{\Theta}_{\alpha_{n-1}, \alpha_{n+1}}^{j_n, j_{n+1}}$ to a matrix and perform an SVD yielding

$$\tilde{\Theta}_{(j_n\alpha_{n-1}), (j_{n+1}\alpha_{n+1})} = \sum_{\alpha_n} U_{\alpha_{n-1}, \alpha_n}^{j_n} S_{\alpha_n, \alpha_n} (V^\dagger)_{\alpha_n, \alpha_{n+1}}^{j_{n+1}}. \quad (115)$$

This causes the bond dimension to increase $D \rightarrow dD$, which must be truncated by keeping only the D largest singular values of S .

4. **Update environments:** The last step depends on which direction one is iterating through the chain. Here the left- and right-normalization of U and V^\dagger is used to update the environments.

Going right: Update the left environment

$$\tilde{L}_{\alpha_n, \beta_n, \alpha'_n}^{[n]} = \sum_{\substack{j_n, j'_n \\ \alpha_{n-1}, \beta_{n-1}, \alpha'_{n-1}}} L_{\alpha_{n-1}, \beta_{n-1}, \alpha'_{n-1}}^{[n-1]} U_{\alpha_{n-1}, \alpha_n}^{j_n} W_{\beta_{n-1}, \beta_n}^{[n]j_n, j'_n} U_{\alpha'_{n-1}, \alpha'_n}^{j'_n*}, \quad (116)$$

and build the matrix of the right site

$$\tilde{M}_{\alpha_n, \alpha_{n+1}}^{[n+1]j_{n+1}} = \sum_{\alpha_n} S_{\alpha_n, \alpha_n} (V^\dagger)_{\alpha_n, \alpha_{n+1}}^{j_{n+1}}. \quad (117)$$

Going left: Update the right environment

$$\tilde{R}_{\alpha_n, \beta_n, \alpha'_n}^{[n+1]} = \sum_{\substack{j_{n+1}, j'_{n+1} \\ \alpha_{n+1}, \beta_{n+1}, \alpha'_{n+1}}} R_{\alpha_{n+1}, \beta_{n+1}, \alpha'_{n+1}}^{[n+2]} \left(V^\dagger \right)_{\alpha_n, \alpha_{n+1}}^{j_{n+1}} W_{\beta_n, \beta_{n+1}}^{[n+1]j_{n+1}, j'_{n+1}} \left(V^\dagger \right)_{\alpha_n, \alpha_{n+1}}^{j_{n+1}*}, \quad (118)$$

and build the matrix of the left site

$$\tilde{M}_{\alpha_{n-1}, \alpha_n}^{[n]j_n} = \sum_{\alpha_n} U_{\alpha_{n-1}, \alpha_n}^{j_n} S_{\alpha_n, \alpha_n}. \quad (119)$$

This concludes the two-site update sequence, which can be seen illustrated in figure 26. After performing the sequence, one can move *one site* to either left or right, depending on which direction one is iterating.

Some comments regarding this sequence are in order: The matrices of the eigenvalue problem has dimensions $(d^2 D^2 \times d^2 D^2)$, which is generally too much for exact diagonalization, however, since only the lowest eigenvalue is of interest, one can use an iterative eigensolver. Furthermore, if the MPS is not in the proper mixed-canonical form, the eigenvalue problem turns into a generalized eigenvalue problem, which can be numerically quite demanding. Thus, updating the left and right environments in step (4) is necessary, since it leads to great simplifications in step (2). Lastly, one could also consider just a single site when updating, however, this method is very prone to getting stuck [42]. By updating two sites at once, one actually optimized the bond between them. Hence, after updating the two sites *one must only iterate a single site* in order to optimize the next bond. Optimization is done with regards to the current configuration, whereby it depends on previous updates. To compensate for this one must sweep through the entire system multiple times, which leads to the following algorithm for iterative ground state search:

Iterative ground state search algorithm

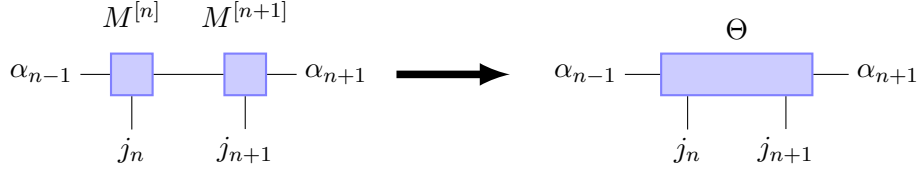
Initialize: Start from some right-normalized initial guess for $|\psi\rangle$. Using this calculate the tensor $R^{[i]}$ iteratively for all sites starting at the right.

Right sweep: Starting from site $n = 1$ perform a *two-site update*, where one updates the *left environment* as described in the sequence. Afterwards n is iterated by 1, $n \rightarrow n + 1$, and the process is repeated until $n = N - 1$ is reached.

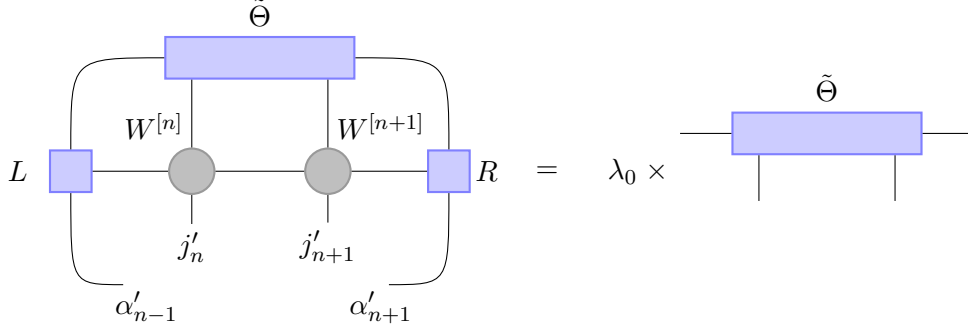
Left sweep: Starting from site $n = N - 1$ perform a *two-site update*, where one updates the *right environment* as described in the sequence. Afterwards n is iterated by 1, $n \rightarrow n - 1$, and the process is repeated until $n = 1$ is reached.

Repeat sweeps: Continue by alternating between left and right sweeps through the system until convergence is achieved.

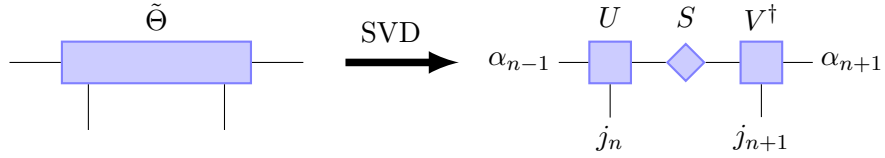
(1) Merge:



(2) Solve eigenproblem:



(3) Unmerge:



(4) Update environments:

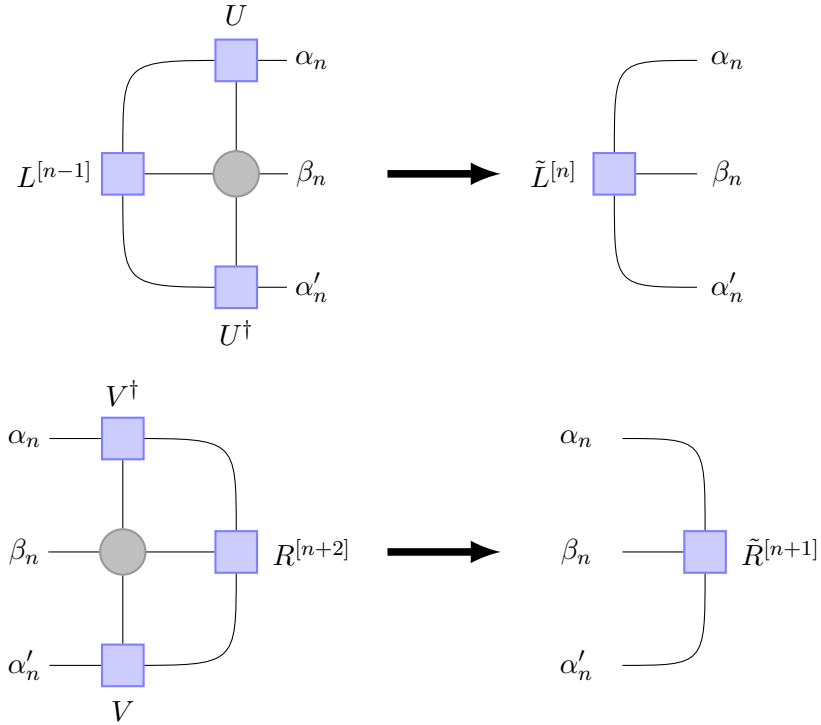


Figure 20: A pictorial representation of the two-site update sequence for iterative ground state search. In step (4) the direction one is iterating determines whether to update the left or right environment.

7 Performing Time Evolution on an MPS

Several algorithms deal with time evolving MPS's, however, as many of them are very similar only a single one, tDMRG, will be treated here.

Consider the time evolution of a quantum state

$$|\psi(t)\rangle = \hat{U}(t) |\psi(0)\rangle, \quad (120)$$

where $\hat{U}(t) = e^{-i\hat{H}t}$ is the time evolution operator. Simply writing this operator as an MPO and applying it is not straightforward, however, if the Hamiltonian contains only nearest-neighbour interactions this can be achieved. Assume the Hamiltonian is a sum of two-site operators of the form $\hat{H} = \sum_n \hat{h}^{[n,n+1]}$. One can decompose this into a sum over even and odd bonds

$$\hat{H} = \hat{H}_{\text{odd}} + \hat{H}_{\text{even}} = \sum_{n \text{ odd}} \hat{h}^{[n,n+1]} + \sum_{n \text{ even}} \hat{h}^{[n,n+1]}. \quad (121)$$

From here one can discretize the time into small time slices δt and perform a Trotter-Suzuki decomposition [36]

$$e^{-i\hat{H}\delta t} = e^{-i\hat{H}_{\text{odd}}\delta t} e^{-i\hat{H}_{\text{even}}\delta t} + O(\delta t^2), \quad (122)$$

where the error is due to the noncommutativity of the bond Hamiltonians $[\hat{h}_{\text{odd}}^{[n,n+1]}, \hat{h}_{\text{even}}^{[n,n+1]}] \neq 0$. Using this the time evolution operator can be written as the product

$$\hat{U}(\delta t) \approx \left(\prod_{n \text{ odd}} \hat{U}^{[n,n+1]}(\delta t) \right) \left(\prod_{n \text{ even}} \hat{U}^{[n,n+1]}(\delta t) \right), \quad (123)$$

where

$$\hat{U}^{[n,n+1]}(\delta t) = e^{-i\hat{h}^{[n,n+1]}\delta t}. \quad (124)$$

Each of these bond evolutions can be written in the form

$$U^{(j_n j_{n+1}), (j'_n j'_{n+1})} = \langle j_n j_{n+1} | e^{-i\hat{h}^{[n,n+1]}\delta t} | j'_n j'_{n+1} \rangle, \quad (125)$$

whereby it can be represented as an MPO. Thus, one has one MPO performing an infinitesimal time step on the odd bonds and another MPO doing the same for even bonds, which can be seen in figure 27.

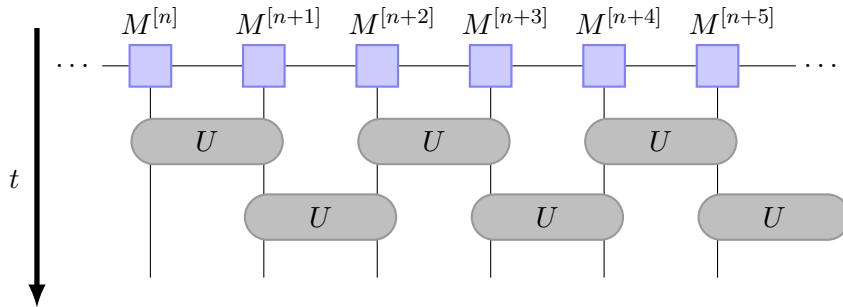


Figure 21: Approximation of each time step δt using a Trotter-Suzuki decomposition, such that the time evolution operator is expressed as a product of unitary two-site operators.

Consider an MPS in the mixed-canonical form

$$|\psi\rangle = \sum_{\mathbf{j}} A^{j_1} \dots A^{j_{n-1}} M^{j_n} S M^{j_{n+1}} B^{j_{n+2}} \dots B^{j_N} |\mathbf{j}\rangle \quad (126)$$

where the matrices M can have any form. Starting from this state the tDMRG update sequence looks as follows:

Infinitesimal time-step update for tDMRG

1. **Merge:** Contract the two matrices $M^{[n]}$ and $M^{[n+1]}$ over the bond α_n creating a two-site tensor

$$\Phi_{\alpha_{n-1}, \alpha_{n+1}}^{j_n, j_{n+1}} = \sum_{\alpha_n} M_{\alpha_{n-1}, \alpha_n}^{[n]j_n} S_{\alpha_n, \alpha_n} M_{\alpha_n, \alpha_{n+1}}^{[n+1]j_{n+1}} \quad (127)$$

2. **Apply unitary:** The two-site time evolution operator, $U^{j_n j_{n+1}, j'_n j'_{n+1}}$, can now easily be applied to the two-site tensor

$$\tilde{\Phi}_{\alpha_{n-1}, \alpha_{n+1}}^{j_n, j_{n+1}} = \sum_{j'_n, j'_{n+1}} U^{j_n j_{n+1}, j'_n j'_{n+1}} \Phi_{\alpha_{n-1}, \alpha_{n+1}}^{j'_n, j'_{n+1}}. \quad (128)$$

3. **Unmerge:** Reshape the updated $\tilde{\Phi}_{\alpha_{n-1}, \alpha_{n+1}}^{j'_n, j'_{n+1}}$ to a matrix and perform an SVD yielding

$$\tilde{\Phi}_{(j_n \alpha_{n-1}), (j_{n+1} \alpha_{n+1})} = \sum_{\alpha_n} U_{\alpha_{n-1}, \alpha_n}^{j_n} S_{\alpha_n, \alpha_n} (V^\dagger)_{\alpha_n, \alpha_{n+1}}^{j_{n+1}}. \quad (129)$$

This causes the bond dimension to increase $D \rightarrow dD$, which must be truncated by keeping only the D largest singular values of S .

4. **Progress:** Next, $e^{-i\hat{h}^{[n, n+1]} \delta t}$ must be applied to the next pair of sites, thus one has to shift by two sites to continue.

First, keep $U_{\alpha_{n-1}, \alpha_n}^{j_n} = A_{\alpha_{n-1}, \alpha_n}^{[n]j_n}$ and form the shifted Φ as

$$\Phi_{\alpha_n, \alpha_{n+2}}^{j_{n+1}, j_{n+2}} = \sum_{\alpha_{n+1}} S_{\alpha_n, \alpha_{n+1}} (V^\dagger)_{\alpha_{n+1}, \alpha_{n+1}}^{j_{n+1}} B_{\alpha_{n+1}, \alpha_{n+1}}^{j_{n+2}}, \quad (130)$$

This is just a shift by one site, however, by reshaping and performing a second SVD one can form the Φ shifted by two sites

$$\Phi_{\alpha_{n+1}, \alpha_{n+3}}^{j_{n+2}, j_{n+3}} = \sum_{\alpha_{n+2}} S_{\alpha_{n+1}, \alpha_{n+2}} (V^\dagger)_{\alpha_{n+2}, \alpha_{n+2}}^{j_{n+2}} B_{\alpha_{n+2}, \alpha_{n+3}}^{j_{n+3}}, \quad (131)$$

where $U_{\alpha_n, \alpha_{n+1}}^{j_{n+1}} = A_{\alpha_n, \alpha_{n+1}}^{[n+1]j_{n+1}}$, thus retaining the canonical form.

After each SVD the matrices are truncated down to dimension D , however, the second SVD causes no loss of information, as the Schmidt rank of S will be at most D .

From here one can apply the unitary $U^{[n+2, n+3]}$ to the new two-site tensor and proceed with the next infinitesimal time-evolution step all while remaining in the canonical form [?]. This sequence is illustrated in figure 28.

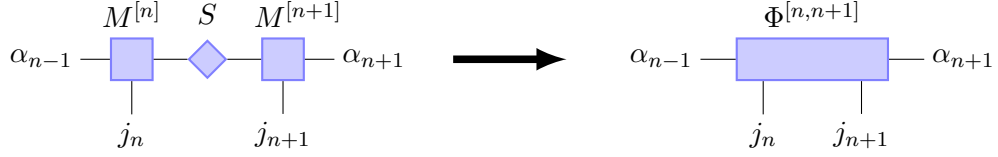
8 Numerical Computations of Bose-Hubbard Model

For the numerical computations the *density-matrix renormalization group* (DMRG) method is used. In this method a general state is represented as a product of matrices, otherwise known as a *matrix product state* or MPS. This is achieved through a series of Schmidt decompositions, where in each iteration the eigenstates of the decomposition contributing the least to the state are truncated. Hence, matrix product states avoids the otherwise exponential growth of the Hilbert space with increasing particle number [32].

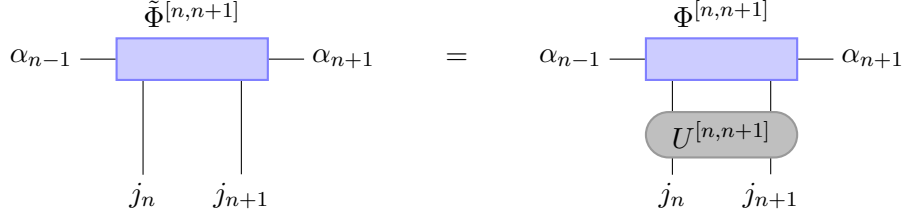
Another important feature of matrix product states is how they follow an area law, meaning the scaling of entropy across the boundary of two regions of the system is linear [7]. Thus, in one dimension the entropy across a cut of the system is constant. This is essential for the success of the MPS representation, as one only has to consider a tiny portion of the full Hilbert space when performing a ground state search.

Hence, the DMRG algorithm, which iteratively performs ground state searches while sweeping over physical sites, is capable of accurately describing one dimensional many-body systems [22, 21].

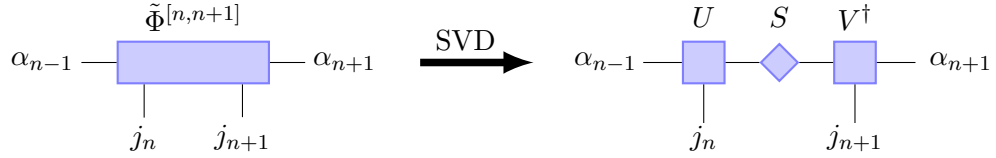
(1) Merge:



(2) Apply unitary:



(3) Unmerge:



(4) Progress:

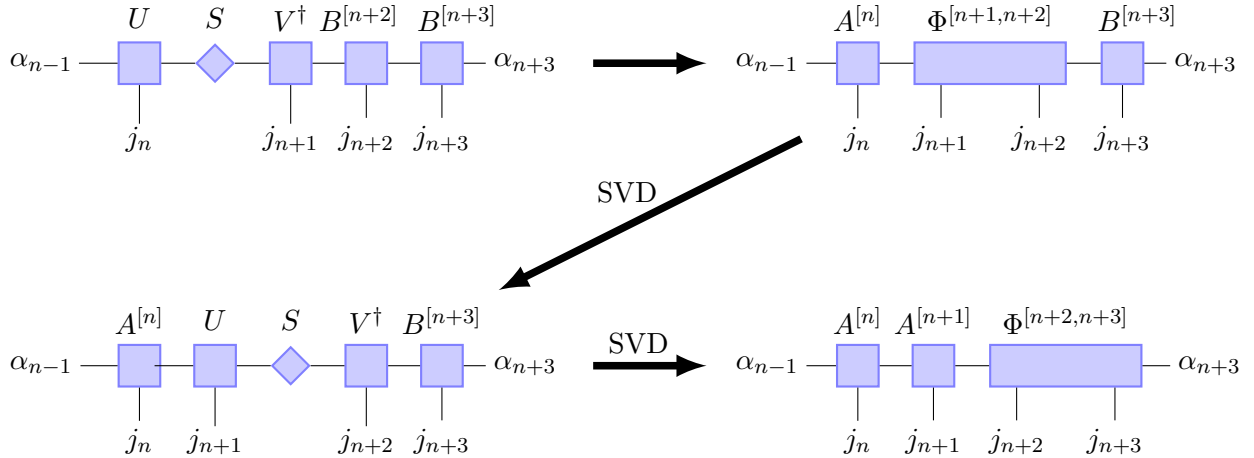


Figure 22: A pictorial representation of the infinitesimal time-step update for tDMRG. After the sequence has been performed, the next unitary $U^{[n+2,n+3]}$ can be applied to the new two-site tensor.

8.1 Correlation Functions

Examining the correlation functions of the Bose-Hubbard model is of special interest, since it relates to the interference pattern of particles measured when performing experiments [17], as well as the Superfluid-Mott Insulator phase transition [22].

Although the MPS formalism excels at describing one dimensional systems, it struggles with describing

the SF correlations, due to how it is constructed. Consider an MPS of the form

$$|\psi\rangle = \sum_{\{j\}} \left(\prod_{n \in \mathbb{Z}} M^{j_n} \right) |\{j\}\rangle, \quad (132)$$

where the M 's are the matrices of the MPS, and j_n are the physical indices. The *transfer operator* is defined as

$$\hat{E}^{[n]} = \sum_{\alpha_{n-1}, \alpha'_{n-1}} \sum_{\alpha_n, \alpha'_n} \left(\sum_{j_n} M^{[n]j_n*} \otimes M^{[n]j_n} \right)_{(\alpha_{n-1}\alpha'_{n-1}), (\alpha_n\alpha'_n)} (|\alpha_{n-1}\rangle \langle \alpha'_{n-1}|) (|\alpha_n\rangle \langle \alpha'_n|), \quad (133)$$

where the expression in the brackets is the matrix elements of the operator, and α_n are the physical indices of the matrices. The transfer operator is essentially a complete, positive map from operators defined on a block of the lattice of length $n-1$ to a block of length n , such that

$$\{|\alpha_{n-1}\rangle \langle \alpha'_{n-1}|\} \rightarrow \{|\alpha_n\rangle \langle \alpha'_n|\}. \quad (134)$$

One important property of the transfer operator, or transfer matrix, is that all eigenvalues $|\lambda_k| \leq 1$ [32].

Generalizing the transfer operator to contraction with an operator \hat{O} gives

$$E_O^{[n]} = \sum_{j_n, j'_n} O^{j_n, j'_n} M^{[n]j_n*} \otimes M^{[n]j'_n}. \quad (135)$$

Using this one can write the correlation function of two general operators on sites i and j as

$$\begin{aligned} \langle \psi | \hat{O}^{[i]} \hat{O}^{[j]} | \psi \rangle &= \text{Tr} E^{[1]} \dots E^{[i-1]} E_O^{[i]} E^{[i+1]} \dots E^{[j-1]} E_O^{[j]} E^{[j+1]} \dots E^{[L]} \\ &= \text{Tr} E_O^{[i]} E^{[j-i-1]} E_O^{[j]} E^{[L-j+i-1]} \\ &= \sum_{l, k} \langle l | E_O^{[i]} | k \rangle \lambda_k^{j-i-1} \langle k | E_O^{[j]} | l \rangle \lambda_l^{L-j+i-1} \\ &= \sum_k \langle 1 | E_O^{[i]} | k \rangle \lambda_k^{j-i-1} \langle k | E_O^{[j]} | 1 \rangle \quad (L \rightarrow \infty) \end{aligned} \quad (136)$$

where λ is the eigenvalues of the transfer matrix, and L is the length of the system. Since $|\lambda_k| \leq 1$, only the leading eigenvalue $\lambda_1 = 1$ remains as $L \rightarrow \infty$. Defining the distance between two sites as $r = |j - i - 1|$ and the correlation decay, or correlation length, as $\xi_k = -1/\ln \lambda_k$, allows one to write the correlation function as

$$\frac{\langle \psi | \hat{O}^{[i]} \hat{O}^{[j]} | \psi \rangle}{\langle \psi | \psi \rangle} = c_1 + \sum_{k=2} c_k e^{-r/\xi_k}, \quad (137)$$

where $c_k = \langle 1 | E_O^{[i]} | k \rangle \langle k | E_O^{[j]} | 1 \rangle$. [32]

Equation (69) shows that the correlation function is a linear combination of exponential functions. Since this was derived without any assumptions regarding neither the MPS nor the operators, this results can be considered general. Thus, any finite-dimensional MPS will only be able to approximate the true correlation of a system.

This is the cause of the difficulty of describing the Superfluid single-particle correlations. While single-particle correlations decay exponentially for the Mott-Insulator, it decays following a power-law for Superfluids

$$\langle \hat{a}_i^\dagger \hat{a}_j \rangle \sim |i - j|^{-K_b/2}, \quad (138)$$

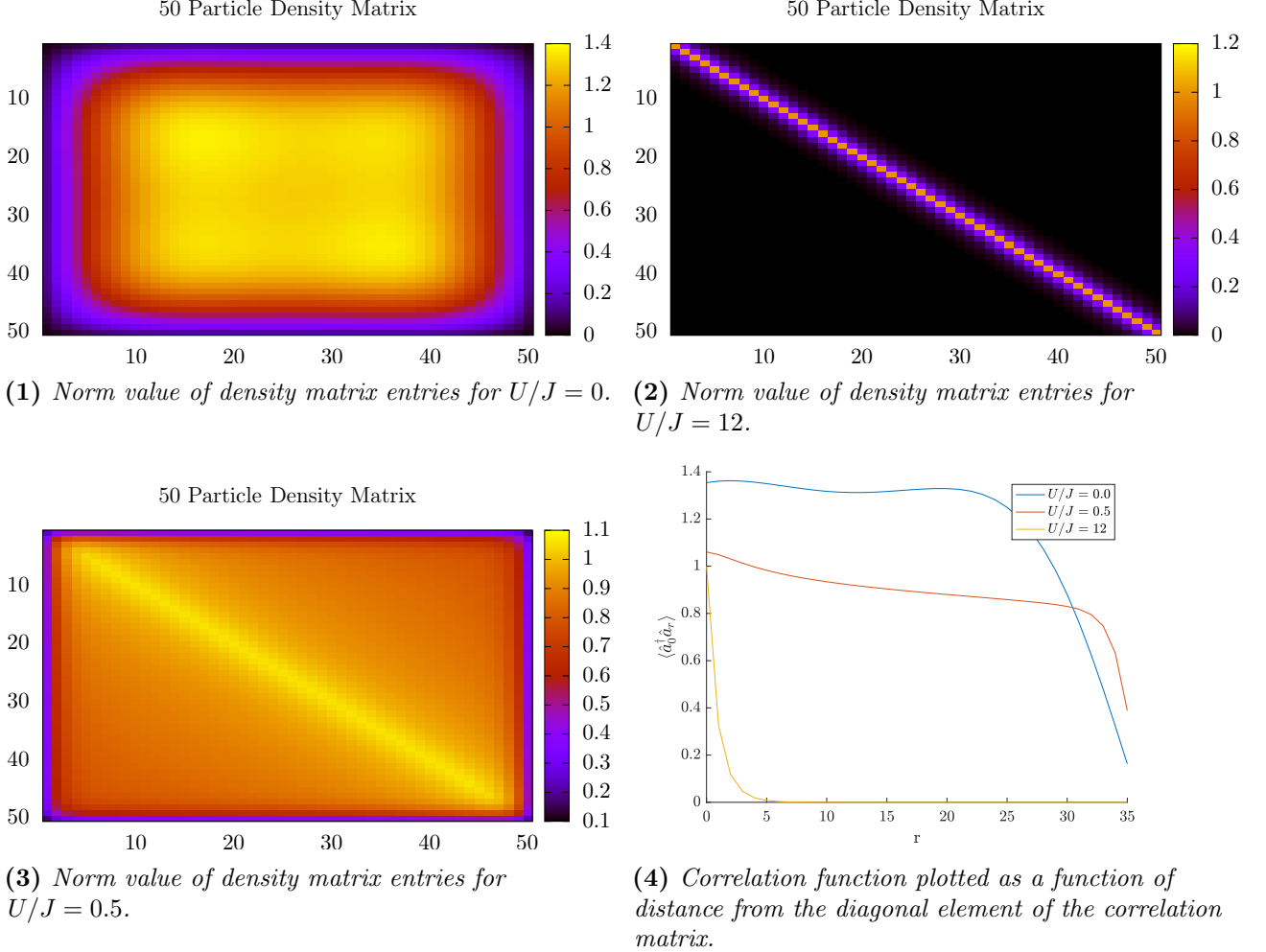
where K_b is the Tomonaga-Luttinger parameter [8]. For short distances equation (69) is able to accurately approximate a power-law, however, as distances grow larger, only the slowest exponential decay will survive. Hence, the correlation turns into a pure exponential decay with $\xi = -1/\ln \lambda$, where λ is the largest eigenvalue of \hat{E} contributing to the correlation.

Visualization of density matrices

One way of displaying the correlations of the system is to plot its density matrix, whose entries are given by

$$\rho_{i,j} = \langle \psi | \hat{a}_i^\dagger \hat{a}_j | \psi \rangle . \quad (139)$$

Figures 8i and 8ii show the density matrix of the 50 particle system plotted for the Superfluid and



the Mott-Insulator limit respectively. In the SF limit long-range correlations are present, which is seen by large off-diagonal elements. However, as discussed previously, in the MPS approximation all correlations decay exponentially, which causes issues when trying to approximate the very long range correlations of the system. This is visualized in figure 8iv where the correlation function is plotted against distance from the diagonal. The Superfluid graph has a hump on it, which is an artefact of the DMRG algorithm. Due to the very long correlation length the graph should be almost flat except for the rapid drop in the end, which is due to the open boundary conditions. The graph in figure 8iv is the result of the DMRG algorithm attempting to express this long range correlation as a series of exponentials.

In the MI limit no interaction takes place between sites and the correlation length is zero, leading to a correlation matrix consisting only of diagonal elements of equal magnitude. Figure (74) shows some off-diagonal elements of non-zero magnitude, however, this system is not a pure Mott-Insulator, as $U/J = 12$. Nevertheless, the correlations are well described by only a single exponential function, as shown in figure 8iv.

Finally, figure 8iii illustrates the density matrix for a system with $U/J = 0.5$, which is primarily as Superfluid, although, as seen in the figure, has a well defined diagonal. Inspecting its correlation function reveals that the best approximation is by a power-law, thus confirming that the system is indeed a Superfluid.

8.2 Condensate Fraction

Recall the Bose-Hubbard Hamiltonian (equation (37)), which supports the two distinct quantum phases: The Superfluid (SF) phase, $U/J \ll 1$, which exhibit long range correlations, and the Mott-Insulator (MI) phase, $U/J \gg 1$, where no interaction between neighboring sites is present. The SF phase is a Bose-Einstein condensate with special attributes, as it exhibits maximal de-localization of the particles and can be approximated as a coherent state. On the other hand, as the individual sites are completely decoupled in the MI phase, no condensate is present.

According to the Penrose-Onsager criterion, a Bose-Einstein condensate, and by extension the SF state, is present if and only if the largest eigenvalue, λ_1 , of the single-particle density matrix, $\rho^{(1)}$, is macroscopic

$$f_c = \frac{\lambda_1}{N_{\text{particles}}} > 0, \quad (140)$$

where f_c is the *condensate fraction* and $N_{\text{particles}}$ is the total number of particles [26]. Thus, the condensate fraction can be used to determine which state is the dominant of the system, as

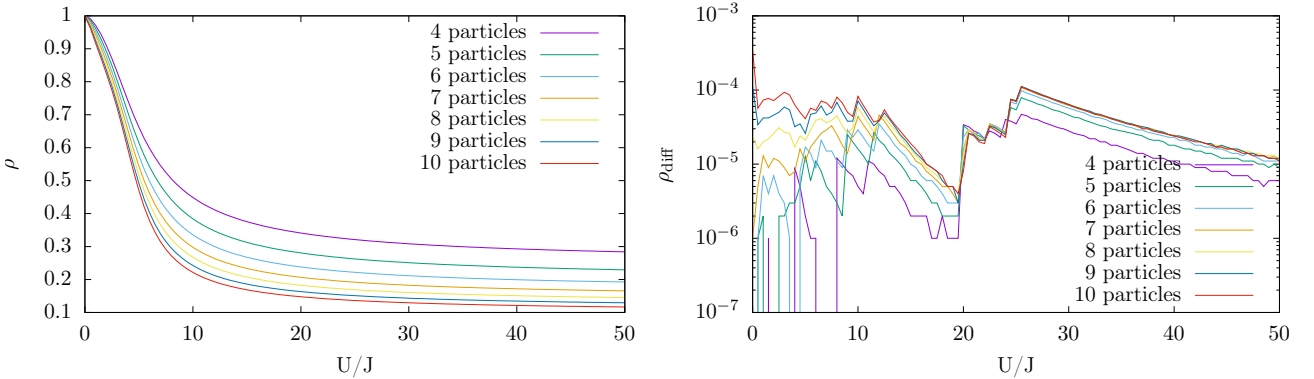
$$\lim_{N_{\text{particles}} \rightarrow \infty} f_c^{\text{SF}} \rightarrow 1 \quad (141)$$

$$\lim_{N_{\text{particles}} \rightarrow \infty} f_c^{\text{MI}} \rightarrow 0, \quad (142)$$

where the filling-fraction $n = N_{\text{particles}}/N_{\text{sites}}$ is held constant. While the limit for the SF phase will be reached for any number of particles, the MI phase will only be reached in the limit $N_{\text{particles}} \rightarrow \infty$.

Calculated condensate fractions

In order to test the precision of MPS formalism and the ITensor library, the condensate fraction was calculated for a series of U/J with varying particle number. For this the library's implementation of the DMRG algorithm was used, where the number of sweeps was selected, such that the bond dimension remained the same when adding more sweeps. In this case 5 sweeps with a maximum bond dimension of 200 was sufficient. All calculations were performed with unit occupancy. These results were then compared to a similar calculation using exact diagonalisation.



(1) Condensate fraction calculated from an MPS after performing 5 DMRG sweeps. (2) Absolute difference between the condensate fraction calculated using MPS and using exact diagonalisation.

Figure 9i shows the condensate fraction for various U/J calculated using MPS. In the limit $U/J = 0$ the condensate fraction is unit as expected, thus confirming that the system is indeed in the SF state. The condensate fraction never reaches zero, as this is only achieved in the thermodynamic limit as described by equation (74). However, the condensate fraction does decrease with increasing particle number as would be expected.

In figure 9ii the results of the MPS calculation is compared to exact diagonalisation by displaying the absolute difference between the two calculations. It is clear that the two approaches yield very similar results for low particle-number.

Attempting to use exact diagonalisation for large particle numbers is futile due to the exponential

scaling of the Hilbert space [40], however, as mentioned, this is not an issue using matrix product states, since low contributing eigenstates are truncated. Figure 10 shows the results of the DMRG

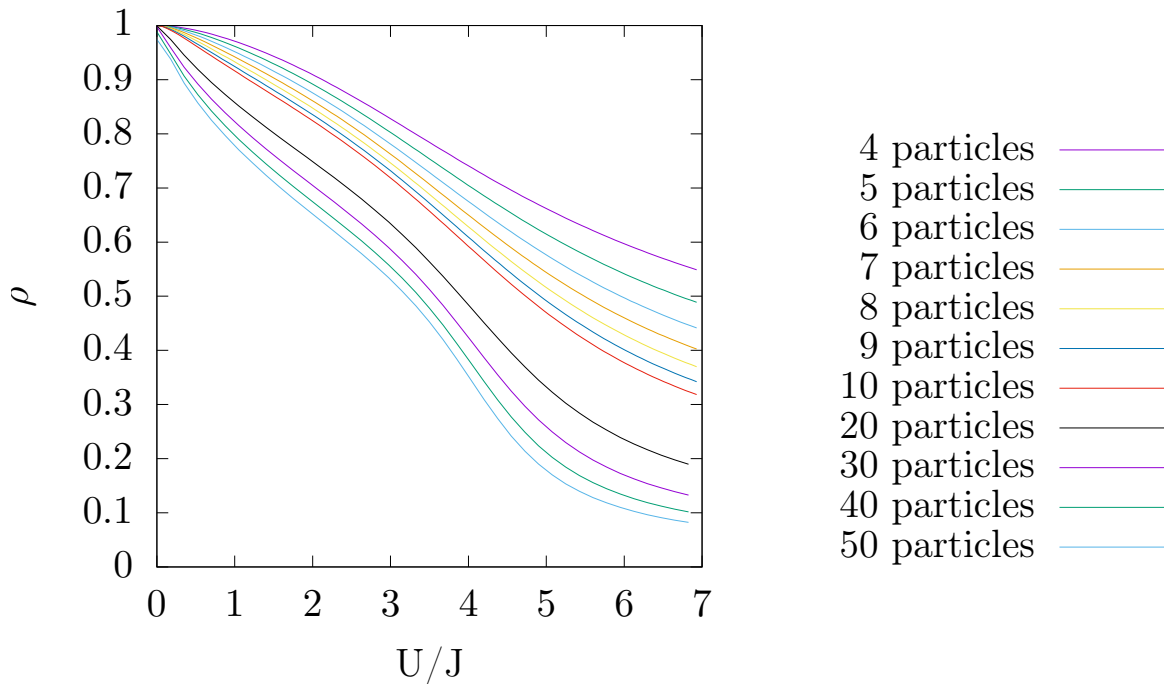


Figure 25: Condensate fraction calculated from an MPS after performing 20 DMRG sweeps.

calculations for up to 50 particles. The condensate fraction behaves as expected in the $U/J \gg 1$ limit, as it tends towards zero for larger particle numbers. Note, that in the SF limit the condensate fraction does not quite reach 1. However, this can be understood when considering figure 8iv, as the DMRG algorithm is unable to describe the very long ranged correlation of the Superfluid. Smaller systems do not have this issue, as boundary effects has an influence on a much larger part of the system. Thus, these systems barely show any sign of long ranged correlations making them easy to approximate.

Some measures can be taken when using the DMRG algorithm in order to minimize errors. Accurately describing long range correlations requires multiple sweeps, as the algorithm iteratively has to approximate an almost constant function with a series of exponentials. Figure 11 displays the condensate fraction in the Superfluid limit as a function of number of sweeps. Clearly, performing more sweeps yields a better result. Furthermore, one observes numerically, that increasing the maximal bond dimension D results in a more accurate long-range representation of correlations. Thus, calculating correlation functions for various values of D is a great way of estimating the convergence of the correlations for a given length scale [32].

In [22] the critical point of phase transition between the Superfluid and Mott-Insulator is determined as $(\frac{U}{J})_{crit} = 3.37$. This is computed by determining the point at which the Luttinger liquid parameter is $K = \frac{1}{2}$. Examining figure 10 one notices a hump on the graph in the vicinity of this critical ratio, however, no clear indication of a phase transition is present. In the thermodynamic limit one would expect the condensate fraction to drop to zero as the critical ratio is reached (as observed in 2D by [35]), although at 50 particles the condensate fraction is only around 0.5. One could extrapolate data from several computations using different particle numbers in order to determine the location of the critical point. However, this would require computations using larger systems in order to minimize the boundary effects.

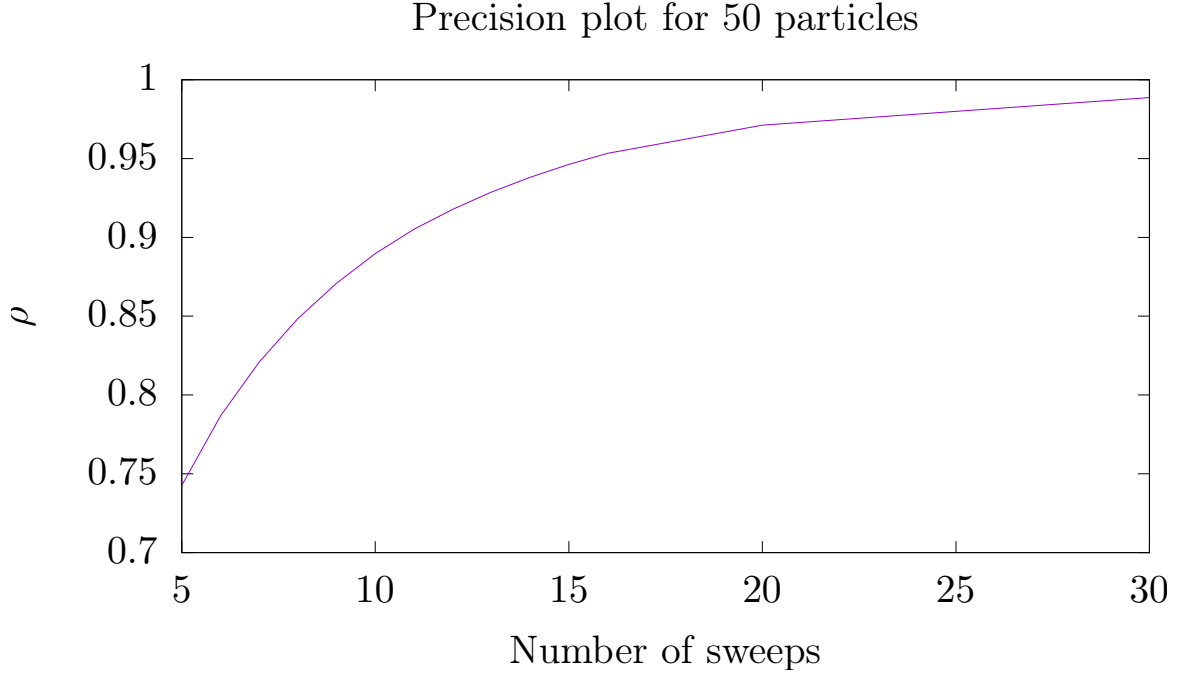


Figure 26: *SF condensate fraction as a function of number of sweeps of the DMRG algorithm. A max bond dimension of 250 was used.*

8.3 Time Evolution

Recall the expression for the superfluid state

$$|\psi\rangle = C \sum_{j=0}^{\infty} \frac{\alpha^j}{\sqrt{j!}} |j\rangle, \quad (143)$$

Recent experimental advances have made it possible to change the Hamiltonian almost instantaneously, allowing for a wide array of new experiments. In the current section, we will look at the case of an SF Hamiltonian,

$$\hat{H} = -J \sum_{\langle i,j \rangle} \hat{a}_i^\dagger \hat{a}_j, \quad (144)$$

present long enough for the condensate to reach the ground state in eq. (75) suddenly being switched to a pure Mott Hamiltonian,

$$\hat{H} = \frac{U}{2} \sum_i \hat{n}_i (\hat{n}_i - 1). \quad (145)$$

In this case, the time evolution of the state can be described by

$$|\psi(t)\rangle = C \sum_{j=0}^{\infty} e^{-it\frac{U}{2}j(j-1)} \frac{\alpha^j}{\sqrt{j!}} |j\rangle, \quad (146)$$

and since $j(j-1)/2 \in \mathbb{N}$ for all j , we will regain the superfluid state at times $p2\pi/U$,

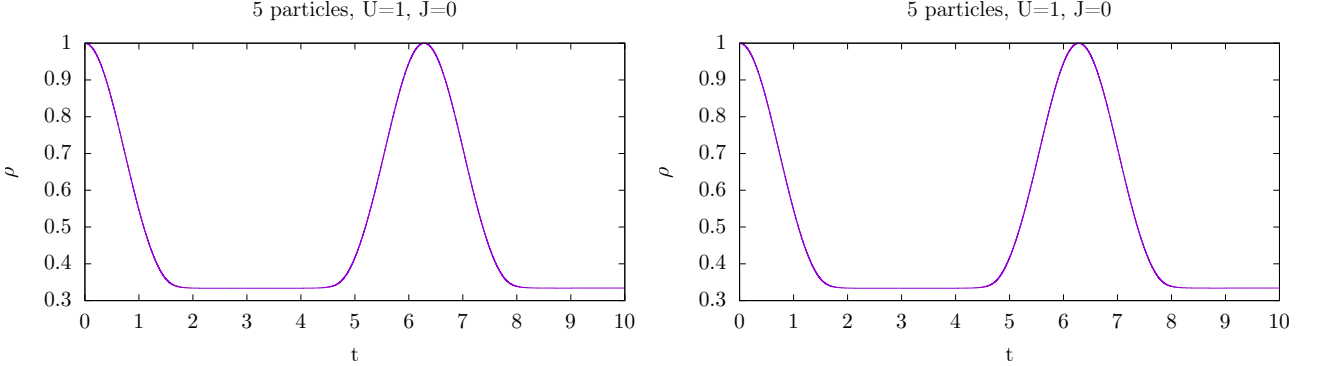
$$|\psi(p2\pi/U)\rangle = |\psi(0)\rangle, \quad (147)$$

for all integers p , whereby we should see a revival of the condensate fraction, as described in ??.

Further testing the applicability of the MPS formalism using ITensor library for time evolution, we evolved an SF state in time first with respect to a pure Mott Hamiltonian ($U = 1$, $J = 0$), then with respect to a slightly different Hamiltonian ($U = 2$, $J = 0.03$).

Time evolution with pure Mott-Insulator Hamiltonian

Mott time evolution on a superfluid state was performed for various unit occupancy systems with complex time steps $t_{1,2} = \frac{1 \pm i}{2}t$ as described in [?], and the condensate fraction was subsequently plotted as a function of time. The initial SF state was found by DMRG with the Hamiltonian in eq. (76). Figure 12i shows time evolution for 5 particles, whereas 12ii is for 50 particles



(1) Condensate fraction calculated from a 5 particle, 5 site MPS after performing 10 DMRG sweeps with an SF Hamiltonian ($J = 1, U = 0$) followed by time evolution with a Mott Hamiltonian ($J = 0, U = 1$).

(2) Condensate fraction calculated from a 50 particle, 50 site MPS after performing 20 DMRG sweeps with an SF Hamiltonian ($J = 1, U = 0$) followed by time evolution with a Mott Hamiltonian ($J = 0, U = 1$).

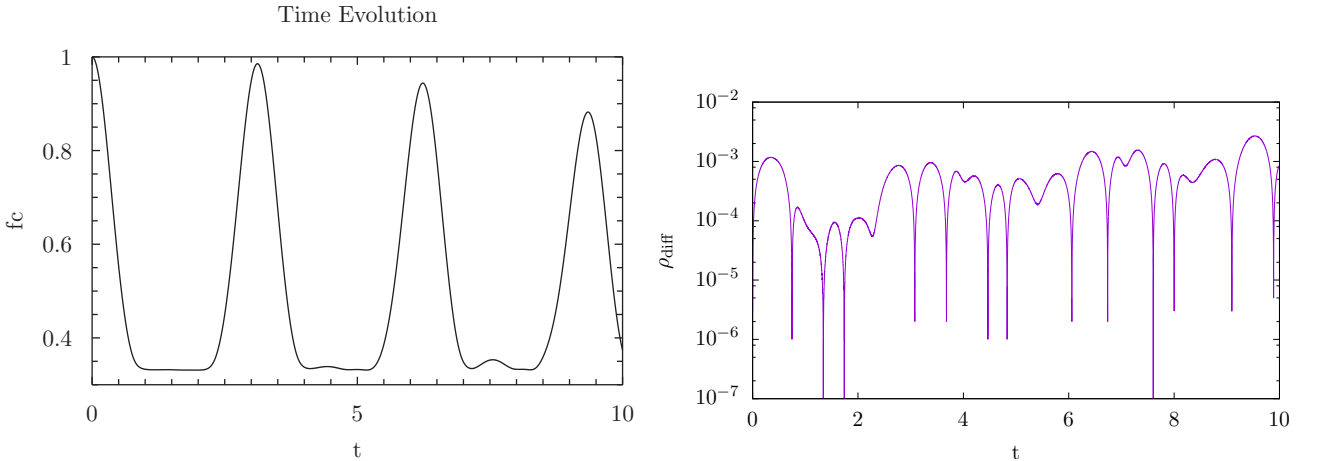
(note: figure not yet made)

with $U = 1$, the period, that is, the time for a full revival of the condensation fraction to occur, can be seen from both figures to be $T = 2\pi$.

WAITING FOR DATA...

Time evolution with mixed Hamiltonian

Performing time evolution on an SF state with a nonzero J , the condensate fraction does not achieve full revival, but instead the phases of the individual particle states interfere with each other more and more for each cycle, leading to a decreasing maximal condensate fraction and some spurious lumps in between the peaks, see fig. 13i. The period of revival is $T = \pi$, since now $U = 2$.



(1) Condensate fraction calculated from a 5 particle, 5 site MPS after performing 10 DMRG sweeps with an SF Hamiltonian ($J = 1, U = 0$) followed by time evolution with a ($J = 0.03, U = 2$) Hamiltonian.

(2) Condensate fraction calculated from a 5 particle, 5 site MPS after performing 10 DMRG sweeps with an SF Hamiltonian ($J = 1, U = 0$) followed by time evolution with a ($J = 0.03, U = 2$) Hamiltonian compared with the same calculations done with exact diagonalization.

In fig. 13ii, the absolute difference between the condensate fraction calculated by MPS formalism and that calculated by exact diagonalization is plotted. The disagreement, while still small at $t = 10$,

is expected to grow larger for larger times, as the MPS does a series of truncations each time the time evolution MPO is applied to avoid increasing the bond dimension of the MPS, whereas exact diagonalization using the Lanczos algorithm detailed in [23] is limited mainly by machine precision.

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