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Abstract

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Reference


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Modulation spectroscopy with ultracold fermions in an optical lattice

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We propose an experimental setup of ultracold fermions in an optical lattice to determine the pairing gap in a superfluid state and the spin ordering in a Mott-insulating state. The idea is to apply a periodic modulation of the lattice potential and to use the thereby induced double occupancy to probe the system. We show by full time-dependent calculation using the adaptive time-dependent density-matrix renormalization-group method that the position of the peak in the spectrum of the induced double occupancy gives the pairing energy in a superfluid and the interaction energy in a Mott insulator, respectively. In the Mott insulator we relate the spectral weight of the peak to the spin ordering at finite temperature using perturbative calculations.

The use of Feshbach resonances [1] and the loading of ultracold atoms into optical lattices [2,3] opened up the possibility to realize strong interactions in quantum gases. These systems provide outstanding control over most of their parameters, which offers the possibility to use them as “quantum simulators” [4], i.e., to investigate unresolved questions from quantum many-body physics in a well-controllable system. One example is the crossover between a Bose-Einstein condensate of bosonic molecules and a BCS-like state of fermionic atoms [5,6]. A major step in the direction of “simulate” phenomena in periodic potentials has been taken by Köhl et al. [7] and very recently by Chin et al. [8] who succeeded in loading ultracold fermions into a three-dimensional optical lattice.

However, the possibilities to probe ultracold gases in optical lattices are still very limited as compared to measurement techniques in condensate-matter systems. On top of this some of the known probes for bosonic gases are not useful in the context of fermionic gases. Thus the recent experimental realization demands new techniques. We propose an experimental setup to measure the pairing energy of a superfluid state and the interaction strength in a Mott-insulating or liquid state to measure the pairing energy of a superfluid state. We further propose how information on the spin ordering could be extracted in the case of repulsive interaction.

In our study we focus on a two-component fermionic gas which is tightly confined to one-dimensional tubes along the x-direction by a strong lattice in the transverse directions. Such a system was realized in the experiment [21]. If an additional periodic potential \( V_x \) is applied along the tubes, the system can be described [12,22] by the Hubbard model

\[
H = -J \sum_{j,\sigma} (c_{j,\sigma}^{\dagger} c_{j+1,\sigma} + H.c.) + U \sum_{j,\sigma} \hat{n}_{j,\uparrow} \hat{n}_{j,\downarrow}.
\]

Here \( \sigma = \{\uparrow, \downarrow\} \) labels the two hyperfine states, \( c_{j,\sigma}^{\dagger} \) and \( c_{j,\sigma} \) are the corresponding creation and annihilation operators, and \( \hat{n}_{j,\sigma} = c_{j,\sigma}^{\dagger} c_{j,\sigma} \) is the number operator on site \( j \). The values of hopping parameter \( J \) and the interaction strength \( U \) depend among other experimental parameters on the lattice potential \( V_x \) [31]. The equilibrium properties of the one-dimensional Hubbard model have been intensively studied by a variety of techniques [23–26], e.g., exact solutions, bosonization, and numerical methods. At low temperature in a system with attractive interaction a superfluid state is formed, whereas for a repulsive interaction the ground state is either a half filled Mott insulator or, away from half filling, a Luttinger liquid. However, the nonequilibrium properties are still a challenge. We use the quasiexact adaptive time-dependent density-matrix renormalization-group method (adaptive t-DMRG) [27–29] and perturbative calculations to determine the time evolution of the system. A periodic modulation of the lattice potential along the tube \( V_x(t) = V_0 [1 + 6 \cos(\omega t)] \), where \( \omega \) is the modulation frequency, can be translated in the lattice description to a periodic modulation of \( J \) and \( U \), i.e., \( J[V_x(t)] \) and \( U[V_x(t)] \). Thus the system is described by an explicitly time-dependent many-body Hamiltonian. In the following, \( J_0 \) and \( U_0 \) will be used for the values \( J[V_0] \) and \( U[V_0] \).

A system subjected to a periodic modulation of the lattice potential will on average absorb energy if the modulation
frequency $\omega$ corresponds approximately to the energy $\Delta E$ needed to create an excitation, i.e., $\hbar \omega = \Delta E$. In the limit of strongly attractive interaction one important class of excitations is the breaking of pairs. In the strongly repulsive limit particle-hole excitations are formed. In both cases these excitations cost an energy of the order of the interaction energy $U_0$. Therefore we expect that the application of a periodic lattice modulation can cause energy absorption, if the frequency corresponds to the excitation energy, i.e., $\hbar \omega = U_0$.

The two cases of attractive and repulsive interaction are related by an electron-hole transformation for one spin species, e.g., $c_{i,j} \rightarrow (-1)^j c_{i,\bar{j}}$, which maps $U \rightarrow -U$ \cite{23,26}. Thus one can restrict the parameters to one sign of the interaction only.

In Fig. 1 the energy absorption for the case of a periodic modulation, both close to resonance $\hbar \omega = U_0$ and off-resonance is shown. In both cases the absorbed energy shows an oscillatory behavior with the frequency corresponding to the modulation frequency. Off-resonance on average no energy is absorbed by the system, whereas at resonance a clear absorption of energy is visible which saturates at longer times.

The energy absorption vs the modulation frequency is shown in Fig. 2. Here the average value at $t_m = 50 \hbar / E_r$ is taken as a measure for the energy absorption \cite{32}. For an initial potential $V_0 = 8 E_r$ a clear peak centered around the resonance frequency $\hbar \omega = U_0$ is seen [Fig. 2(a)]. For a lower initial potential $V_0 = 4 E_r$ [Fig. 2(b)] we see a broadening of the peak and a clear asymmetric form. The maximum of the peak is shifted slightly above the value $\hbar \omega = U_0$. These changes are due to the fact that the distance of the ground-state energy to the center of the next band is not exactly given by the energy $U_0$ but shows corrections in $J_0$.

In the experiment with bosonic atoms \cite{13} the broadening of the characteristic low momentum peak in the time-of-flight images has been used as a measure for the absorbed energy. In the fermionic experiment the observation of the absorbed energy is more difficult. Due to Pauli’s principle for the momentum distribution all momenta up to the Fermi momentum are occupied. Therefore the absorption of energy will only cause a smearing out of the step around the Fermi momentum in time-of-flight images which cannot be used to determine quantitatively the amount of energy absorbed \cite{33}

We propose to use the double occupancy in the lattice, i.e., $D = \langle \sum p_{j,\uparrow} p_{j,\downarrow} \rangle$, as a quantity which can be observed more easily by projecting pairs of atoms onto molecules \cite{20}. Since for strongly attractive (repulsive) interactions the energy absorption corresponds to a breaking of pairs (generation of particle-hole excitations) we expect that in this limit the energy absorption shows up in the expectation value of the double occupancy. In perturbative calculations we find that up to the quadratic response regime the average induced double occupancy and the average absorbed energy are related simply by a factor $U_0$. This result is obtained by expanding $J(t)$ up to first order in $\delta V$, neglecting the time dependence in the interaction, i.e., $U(t) = U_0$, and writing the Hamiltonian as $H(t) = H_0 + g(t) H_K$, where $H_0$ is the Hubbard Hamiltonian at $t = 0$, $H_K$ is the hopping operator, and $g(t) = (d J / d V_0) \Delta V = \delta V \cos(\alpha t)$. The linear response to this time-dependent perturbation gives a purely oscillating contribution to the double occupancy, which averages to zero. Therefore we consider the quadratic response, which is given by

$$D_2(t) = \frac{1}{U_0^2} \int_{-\infty}^{\infty} dt' \int_{-\infty}^{\infty} dt'' g(t') g(t'') \chi_2(t, t', t''),$$

where $\chi_2(t, t', t'')$ is the quadratic susceptibility. Its relevant term, i.e., the one that gives a nontrivial time dependence, can be related to the first-order susceptibility,

$$\chi_2^{\text{rel}}(t, t', t'') = - \theta(t' - t'') \frac{d}{dt} \chi_k(t - t') + \text{(osc. terms)},$$

where $\theta(t)$ is the step function, and $\chi_k(t) = (1/\hbar) \times \langle [H_0(t), H_K(0)] \rangle$. The final expression for the double occupancy is

$$D_2(t) = D_0(1 - \frac{1}{4} \langle g(0) \rangle^2) \text{Im} \chi_k(\omega)/U_0 + \text{(osc. terms)},$$

where $\chi_k(\omega)$ is the Fourier transform of
FIG. 3. (Color online) The position of the peak in the induced double occupancy. Results obtained by a full time-dependent calculation (adaptive t-DMRG) are shown at half filling, \(a_s=9.2\ \text{nm}\) for a strong perturbation \(\delta V=0.2\) (squares), at half filling for a weak perturbation \(\delta V=0.01\) (triangles), and away from half filling for a strong perturbation \(\delta V=0.2\) (circles). Perturbative results (dashed line) are only shown for the temperature \(k_B T=1/\beta=0.1 U_0\), since the results for \(k_B T=0.01 U_0\) and \(k_B T=0.02 U_0\) coincide within the thickness of the line (\(L=6\) and \(N=6\)). A good agreement between all results is found, which makes the position of the peak a good measure of \(U_0/J_0\).

\(\chi_0(t)\). One finds the same linear growth with time for the expectation value of the total energy, without the factor \(1/U_0\). The oscillatory terms, however, are different for both quantities.

In the experimental realization a relatively strong modulation strength \(\delta V\) should be used to get an observable signal. We checked numerically that even for modulation strengths of 20%, the structure of the energy absorption spectrum and the spectrum of the induced double occupancy still contain similar information. In particular we find that the position of the peak agrees for both, whereas deviations occur in the width and amplitude for higher values of the perturbation. Examples for the comparison are shown in Fig. 2.

Since we have seen that the experimentally measurable quantity, the induced double occupancy, can be used as a similar measure as the absorbed energy, we will discuss in the following what kind of information can be extracted from this quantity. Hereby we focus on (i) how to obtain a good measure of the energy necessary to break a pair in the superfluid state or to create a particle-hole excitation in a Mott-insulator or liquid state, and (ii) how to gain information about the spin ordering in a state.

Direct access to the average energy needed to break pairs or to create a particle hole excitation can be obtained from the position of the absorption peak. At zero temperature the dependence of the peak position on the initial interaction strength \(U_0/J_0\) is shown in Fig. 3 for different modulation strengths and fillings (marked by squares, triangles, and circles). The results were obtained performing the full time evolution of the system using the adaptive t-DMRG [34].

The position of the peak depends only on the initial interaction strength and not on the size of the perturbation. For strong interactions the dependence is linear. For small values of the interaction strength \(U_0/J_0\) it deviates from the linear form, since the delocalization of the fermions becomes more important and the simple picture of assuming the state to consist of localized particles breaks down [23,24]. To study the robustness of the peak to the influences of finite temperature we additionally perform a perturbative calculation for small system sizes. In Fig. 3 we show the position of the maximum value of the contributions in quadratic response calculations [35]. The collapse of the data for different temperatures shows that the position is mainly independent of temperature. The good agreement between our results shows that the position is a very robust quantity, which is almost independent of temperature, strength of the modulation, size, and filling of the system.

In experiments an additional harmonic trapping potential is present. We expect that its main influence on the induced double occupancy is a broadening of the width of the peak, as was found in the bosonic case [19]. In particular, the position of the peak will remain unaffected. In the limit of strong interaction this can be understood by the introduction of potential-energy differences between neighboring sites by the trapping potential. This causes a shift of the excitation energy for the pair breaking or the creation of particle-hole pairs, but with a sign depending on the direction of the excitation. Therefore the position of the peak remains unchanged whereas its form can be broadened. In the experiment with bosonic atoms [13] for strong interaction strength the position of the resonance peak in the energy absorption could be determined approximately up to an accuracy of 20 Hz. Note that since the broadening of the peak is proportional to \(J_0\) the accuracy is best for strong interaction. In the fermionic case it is hard to estimate the loss of accuracy by the projection onto molecules, but we expect that the accuracy which can be obtained is of the same order of magnitude.

In a two-component fermionic system with repulsive interactions the probability to create a particle-hole excitation depends strongly on the spin ordering present in the initial state. In a ferromagnetically ordered state with only a few domain walls particle-hole excitations are strongly suppressed by the Pauli principle, whereas in an antiferromagnetically ordered state the creation of particle hole pairs is facilitated. This means that a state which is antiferromagnetically ordered will show more absorption than a paramagnetically or ferromagnetically ordered state. Thus the weight of the ab-
sorption peak gives information about the spin ordering. To be more specific, for large $U_0/J_0$ perturbative calculations yield that the weight is proportional to the nearest-neighbor correlation $\langle n_{i} n_{i+1}\rangle$. This can be seen in Fig. 4 where we compare perturbation results for the weight of the absorption and nearest-neighbor correlations for different temperatures. At large values of $U_0/J_0$ we find very good agreement between the quantities, whereas for smaller interaction the values deviate. A maximum occurs in both quantities at approximately the same finite interaction strength. This is due to two competing processes which destroy the spin order, the delocalization of charge dominating at low $U_0/J_0$, and the spin exchange dominating at large $U_0/J_0$. In the inset in Fig. 4 this competition of the energy scales for the spin ordering, the charge gap, and the spin-exchange coupling are shown. The maximum in the weight shows the crossover between the two energy scales thus giving information about the spin ordering of the system.

To summarize, we have proposed a setup to measure the pair-binding energy, the interaction strength, and the spin ordering in a Fermi gas in an optical lattice. The proposed experiment observes the double occupancy induced by periodic lattice modulations via the creation of molecules. The robustness of the measurement procedure to external influences such as temperature or a harmonic trapping potential makes a precise experimental determination of the pair-binding energy, the interaction strength—thereby the scattering length—and the spin ordering in the optical lattice possible. Since for strong interactions the proposed measurement is based on a local effect, i.e., the creation of particle-hole pairs, we expect it to hold as well in higher dimensions.

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[31] In our simulation we use parameters corresponding to a gas of K40 atoms in the F=9/2, m_f=−9/2, and m_f=−7/2 states, and λ=826 nm.
[32] This is determined by smoothing the oscillations in the curve by convolution with a Gaussian $\exp\left[-r^2/(2\Delta^2)\right]$ with $\Delta T = \pi/\omega$.
[33] A similar smearing is caused by the finite temperature and the trapping potential [30].
[34] We determine the position $a$ by fitting a Gaussian of the form $c \exp\left[-(x-a)^2/\sigma^2\right]$. Note that the fit by a Gaussian function is not very accurate for small values of $U_0/J_0$ due to the asymmetric form of the peak.
[35] We show the position of the maximum value of the $\delta$ peaks, i.e., $1/\langle Z(E_n−E_0)|\langle n|H_K|m\rangle|^2\exp\left[-\beta(E_n−E_0)(1−\exp\left[-\beta(E_n−E_0)\right])\right]$. Here $E_n$ are the eigenvalues of the unperturbed Hamiltonian, $\beta$ gives the inverse temperature.