

Magnetic quantum phase transition of cold atoms in an optical lattice

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We propose a scheme to investigate the magnetic phase transition of cold atoms confined in an optical lattice. We also demonstrate how to get coupled two-leg spin ladders which display a phase transition from a spin liquid to magnetic ordered state in two-dimensional optical lattice. An experimental protocol is further designed for observing this phenomenon.

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I. INTRODUCTION

Cold atoms in optical lattices provide a useful experimental means to investigate quantum many-body systems due to the advantage of cleanness and controllability. Extensive works have been done in the past few years, such as implementing quantum phase transitions from a superfluid to a Mott insulator [1,2], simulating high- T_c superconductivity [3], and realizing various Hubbard models and spin models [4,5]. By adjusting the amplitudes and propagation directions of laser beams, a variety of geometries of optical lattice are generated [6,7]. This technique offers the possibility for constructing spin systems including spin chains [8–10], kagome lattices [11], and spin ladders [8,12,13]. In experiments, except for cold Bose atoms, Fermi atoms in optical lattices have also been realized recently, such as ^{40}K in one-dimensional (1D) [14] and 3D [15] lattices, ^6Li in 3D lattices [16], and a variety of interesting phenomena—for example, Bloch oscillations, band insulators, and superfluidity—were reported.

It was known that spin ladders may provide a transition from 1D chains to 2D lattices [17]. A class of cuprates can be described by antiferromagnetic spin ladders with spin 1/2 [18–21]. Applying a magnetic field or pressure, these cuprates show a phase transition from a spin liquid state to several ordered magnetic states. Strongly correlated cold atoms in optical lattices provide a way to observe this quantum phase transition.

In this paper, we propose an optical lattice setup to produce 2D coupled spin ladders to illustrate a quantum phase transition, from spin-dimerized phase to magnetically ordered phase, and further discuss experimental conditions which can be realized by adjusting the lattice parameters. Compared to cuprates, optical lattices are clean and easily controllable. Manipulating the amplitudes and wave vectors of laser beams, the coupling between spins can be adjusted in a wide range.

II. OPTICAL LATTICES AND SPIN LADDERS

A 2D superlattice is formed by superimposing a standing wave along the x direction with twice the period of a 2D optical lattice, which was generated by two perpendicular

standing waves, as illustrated in Figs. 1(a) and 1(b). Adjusting the intensities of the three standing waves, the tunneling and potential barriers along the x and y direction can be well controlled. For sufficiently large intensities of the laser, the on-site interaction strength is much larger than the kinetic energy so that the lattice-atom system can be in the Mott insulating phase at a commensurate filling. We will consider an equal-mixing two-component fermionic system and assume that the lattice has been loaded with one atom per lattice site. This is possible by using the coherent filtering scheme proposed in Ref. [22].

In general, the Hamiltonian of interacting cold atom gas in an optical lattice is written as

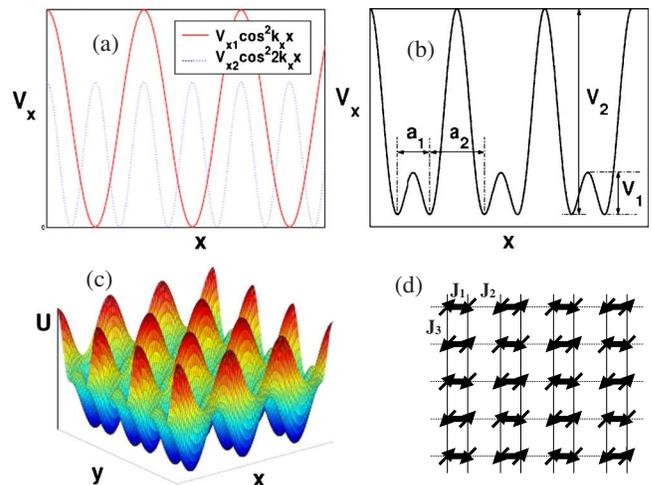


FIG. 1. (Color online) (a) Two selected optical potentials along the x direction, where the potential V_{x2} (dotted line) has twice the period of V_{x1} (solid line). (b) Total potential in the x axis, where a_1 , a_2 , V_1 , and V_2 are the widths and heights of barriers of intradimers and interdimer, respectively. (c) Landscape of potential in the x - y plane. (d) Geometry of the 2D coupled spin ladders, where J_1 , J_2 , and J_3 are intradimer, interdimer, and interladder spin-spin couplings, respectively.

$$H = \sum_{\sigma} \int d\mathbf{r} \hat{\Psi}_{\sigma}^{\dagger}(\mathbf{r}) \left[-\frac{\hbar^2}{2m} \nabla^2 + U(\mathbf{r}) \right] \hat{\Psi}_{\sigma}(\mathbf{r}) + \sum_{\sigma\sigma'} \frac{g}{2} \int d\mathbf{r} \hat{\Psi}_{\sigma}^{\dagger}(\mathbf{r}) \hat{\Psi}_{\sigma'}^{\dagger}(\mathbf{r}) \hat{\Psi}_{\sigma'}(\mathbf{r}) \hat{\Psi}_{\sigma}(\mathbf{r}), \quad (1)$$

where $g=4\pi a_s \hbar/m$ and a_s is the s -wave scattering length. The potential of optical lattice has the form

$$U(\mathbf{r}) = V_{x1} \cos^2(k_x x) + V_{x2} \cos^2(2k_x x) + V_y \cos^2(k_y y) + (m\omega_z^2 z^2)/2 + V_{x1}^2/(16V_{x2}) - V_{x1}/2, \quad (2)$$

where V_{x1} and V_{x2} are the barrier heights of the two standing waves along the x direction, V_y is that along the y direction, k_x and k_y are the two components of wave vector along the x and y directions, and ω_z is the harmonic frequency of the potential in the z direction. The last two terms make the potential value zero in the bottom of every minitrap. To simulate a 2D coupled two-leg spin ladder, $4V_{x2} > V_{x1}$ is necessary. The heights of the two potential barriers in the x direction are $V_1=(4V_{x2}-V_{x1})^2/(16V_{x2})$ and $V_2=(4V_{x2}+V_{x1})^2/(16V_{x2})$. The widths of the two barriers in the x direction and the one in the y direction are $a_1=\arccos(V_{x1}/4V_{x2})/k_x$, $a_2=[\pi-\arccos(V_{x1}/4V_{x2})]/k_x$, and $b=\pi/k_y$. The geometry of the optical lattice is displayed in Fig. 1(b).

In the bottom of the trap, we take the harmonic approximation and the frequencies in the x and y directions are $\omega_x^2=(16V_{x2}^2-V_{x1}^2)k_x^2/(2mV_{x2})$ and $\omega_y^2=2V_y k_y^2/m$, respectively. When the on-site interaction and thermal fluctuation are much smaller than the excited energy of the second band, all atoms are located in the lowest band. The field operators can be expanded as $\hat{\Psi}_{\sigma}(\mathbf{r})=\sum_{i,j} c_{i,j,\sigma} w(\mathbf{r}-\mathbf{r}_{i,j})$, and $w(\mathbf{r}-\mathbf{r}_{i,j})$ is the ground-state function of the harmonic oscillators, $w(\mathbf{r})=[m\bar{\omega}/(\pi\hbar)]^{3/4} e^{m(\omega_x x^2 + \omega_y y^2 + \omega_z z^2)/(2\hbar)}$. $c_{i,j,\sigma}$ are the annihilation operators for spin- σ atoms localized at the site labeled by (i,j) , and $\bar{\omega}$ represents the geometric mean of frequencies, $\bar{\omega}=(\omega_x \omega_y \omega_z)^{1/3}$. Substituting the field operator into the Hamiltonian and integrating, we obtain an equivalent 2D Hubbard model

$$H = - \sum_{i,j,\sigma} (t_1 c_{2i,j,\sigma}^{\dagger} c_{2i+1,j,\sigma} + t_2 c_{2i+1,j,\sigma}^{\dagger} c_{2i+2,j,\sigma} + \text{H.c.}) - \sum_{i,j,\sigma} (t_3 c_{i,j,\sigma}^{\dagger} c_{i+1,j,\sigma} + \text{H.c.}) + U \sum_{i,j,\sigma} n_{i,j,\sigma} n_{i,j,\bar{\sigma}}, \quad (3)$$

where $\sigma=\pm 1/2$ represents the two states of cold atoms and $\bar{\sigma}=-\sigma$. The hopping and repulsive parameters are written as $t_1=e^{-m\omega_x a_1^2/4\hbar} (t_0 + \frac{m\omega_x^2 a_1^2}{8} + \frac{V_{x1}}{2} e^{-\hbar k_x^2/m\omega_x})$, $t_2=e^{-m\omega_x a_2^2/4\hbar} (t_0 + \frac{m\omega_x^2 a_2^2}{8} - \frac{V_{x1}}{2} e^{-\hbar k_x^2/m\omega_x})$, and $t_3=e^{-m\omega_y b^2/4\hbar} (t_0 + \frac{m\omega_y^2 b^2}{8} - \frac{V_{x1}}{2} e^{-\hbar k_x^2/m\omega_x})$, where $t_0=-\frac{\hbar}{4}(\omega_x + \omega_y + 2\omega_z) - \frac{V_{x2}}{2}(1 + e^{-4\hbar k_x^2/m\omega_x}) - \frac{V_y}{2}(1 + e^{-\hbar k_y^2/m\omega_y}) - \frac{V_{x1}^2}{16V_{x2}}$ and $U=\sqrt{\frac{2}{\pi}} \hbar \bar{\omega} a_s / \bar{a}_0$, with $\bar{a}_0=\sqrt{\hbar/(m\bar{\omega})}$.

In the regime of strong coupling, $U \gg t$, the half-filled Hubbard model is equivalent to a spin model approximately up to the order of $t_{1,2,3}^2/U$,

$$H = \sum_{i,j} [J_1 \mathbf{S}_{i,j}^l \cdot \mathbf{S}_{i,j}^r + J_2 \mathbf{S}_{i,j}^r \cdot \mathbf{S}_{i+1,j}^l + J_3 (\mathbf{S}_{i,j}^l \cdot \mathbf{S}_{i,j+1}^l + \mathbf{S}_{i,j}^r \cdot \mathbf{S}_{i,j+1}^r)], \quad (4)$$

where $J_m=t_m^2/(4U)$, $m=1,2,3$, and $\mathbf{S}_{i,j}^{l,r}$ are the spin-1/2 operators on the left and right of the (i,j) dimer. This Hamiltonian describes a 2D coupled spin ladder, as displayed in Fig. 1(d).

In real systems, an additional weak isotropic harmonic potential exists over the lattice [2]. This confinement brings an energy offset of each lattice site and leads to an effective local chemical potential. The superfluid-insulator phase diagram is influenced by this potential [1]. In the current case, the contribution of this potential only adds a term in Eq. (4):

$$m\omega_{wh}^2 \sum_{i,j,m} [(i-N_1/2)^2 a^2 + (j-N_2/2)^2 b^2] (\mathbf{S}_{i,j}^m)_z, \quad (5)$$

where $m=l,r$ and ω_{wh} denotes the trapping frequency of the whole harmonic potential. N_1 and N_2 are the number of dimers along the x and y directions, respectively. Similarly, a and b represent the space of dimers. The effect of this whole harmonic potential corresponds to producing an effective magnetic field gradient in the z direction. For a typical optical lattice, this potential is very weak; for example, in Ref. [2], the trapping frequency is about 65 Hz. For a lattice with ~ 100 sites in a single direction, the Zeeman energy for atoms in different sites varies from $\sim 10^{-5}E_r$ to $\sim 10^{-2}E_r$. The magnitude is much smaller than the coupling energy. Thus, this additional harmonic potential is ignored in the following discussion.

III. PHASE TRANSITION

In the following, we consider repulsive spin-1/2 atoms [23,24] and focus on the magnetic properties of the system described in Eq. (4). By adjusting the optical lattice parameters, we can reach the regime that $J_1 > J_2(J_3)$. In this case, the bond operator method [25] is appropriate to study the properties of the system at low temperature. It describes well the dimerized phase and several magnetically ordered phases of dimer-based antiferromagnets. For a dimer of two 1/2 spins, the Hilbert space can be spanned by four states: one spin singlet $|s\rangle=(|\uparrow\downarrow\rangle-|\downarrow\uparrow\rangle)/\sqrt{2}$ and three spin triplets $|t_x\rangle=-(|\uparrow\uparrow\rangle-|\downarrow\downarrow\rangle)/\sqrt{2}$, $|t_y\rangle=i(|\uparrow\uparrow\rangle+|\downarrow\downarrow\rangle)/\sqrt{2}$ and $|t_z\rangle=(|\uparrow\downarrow\rangle+|\downarrow\uparrow\rangle)/\sqrt{2}$. Bond operators s^{\dagger} and $t_{\alpha}^{\dagger}(\alpha=x,y,z)$ are introduced to generate the singlet and triplet states out of the vacuum $|0\rangle$, $|s\rangle=s^{\dagger}|0\rangle$, $|t_x\rangle=t_x^{\dagger}|0\rangle$, $|t_y\rangle=t_y^{\dagger}|0\rangle$, and $|t_z\rangle=t_z^{\dagger}|0\rangle$. The bond operators s and t_{α} are assumed to satisfy the bosonic communication relations with a constraint of single occupancy at each bond, $s_{i,j}^{\dagger} s_{i,j} + t_{i,j}^{\alpha\dagger} t_{i,j}^{\alpha} \equiv 1$.

In terms of these four operators, the two spin operators in a dimer can be expressed as $(S_{i,j}^l)_{\alpha}=(s_{i,j}^{\dagger} t_{i,j}^{\alpha} + t_{i,j}^{\alpha\dagger} s_{i,j}) - i\epsilon_{\alpha\beta\gamma} t_{i,j}^{\beta\dagger} t_{i,j}^{\gamma}$ and $(S_{i,j}^r)_{\alpha}=(s_{i,j}^{\dagger} t_{i,j}^{\alpha} - t_{i,j}^{\alpha\dagger} s_{i,j} - i\epsilon_{\alpha\beta\gamma} t_{i,j}^{\beta\dagger} t_{i,j}^{\gamma})/2$, respectively, where α, β , and γ represent x, y , and z , and ϵ is the fully antisymmetric Kronecker symbol. Repeated indices are summed over. Under the above transformation, the spin Hamiltonian, Eq. (4) can be transferred into a bosonic one. The constraint condition can be realized by introducing the

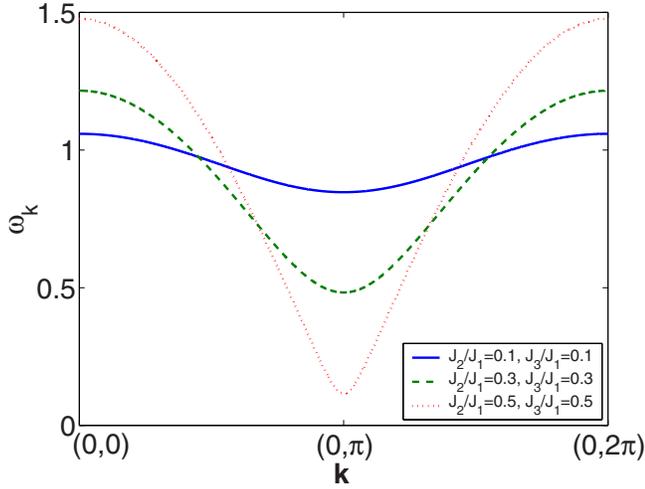


FIG. 2. (Color online) The energy spectrums of different inter-dimer couplings in the spin liquid phase. The solid, dashed, and dotted lines correspond to coupling strength $J_2/J_1=J_3/J_1=0.1, 0.3, 0.5$, respectively. The optical lattice parameters $V_{x1}=5E_r$, $V_{x2}=20E_r$, and $V_y=28E_r$ for $J_2/J_1=J_3/J_1=0.1$; $V_{x1}=16E_r$, $V_{x2}=10E_r$, and $V_y=6E_r$ for $J_2/J_1=J_3/J_1=0.3$; $V_{x1}=20E_r$, $V_{x2}=12E_r$, and $V_y=7E_r$ for $J_2/J_1=J_3/J_1=0.5$. $E_r=\frac{\hbar^2 k_x^2}{2m}$, where m is the mass of the ${}^6\text{Li}$ atom, $a_s=2.41$ nm, $m=9.99\times 10^{-26}$ kg, $k_x=2k_y=5.93$ μm^{-1} , and $\omega_z=6600$ Hz. ω_k is measured in units of J_1 . k_x and k_y are measured in units of $1/a$ and $1/b$, respectively. In the following figures, the atom parameters and laser wavelengths are the same.

Lagrange multipliers $\mu_{i,j}$ in the Hamiltonian, $-\sum \mu_{i,j}(s_{i,j}^\dagger s_{i,j} + t_{i,j}^{\alpha\dagger} t_{i,j}^\alpha - 1)$. Performing the Fourier transformation $t_{i,j}^\alpha = 1/\sqrt{N} \sum_{\mathbf{k}} t_{\mathbf{k}}^\alpha e^{i\mathbf{k}\cdot\mathbf{r}_{i,j}}$, $s_{i,j} = 1/\sqrt{N} \sum_{\mathbf{k}} s_{\mathbf{k}} e^{i\mathbf{k}\cdot\mathbf{r}_{i,j}}$ (N is the number of dimers), and taking $s_{\mathbf{k}} = s_{\mathbf{k}}^\dagger = \tilde{s}$ as that most of the s are condensed in the condition $J_1 \gg J_{2,3}$, we get $H = \sum_{\mathbf{k}} [\Gamma_{\mathbf{k}} t_{\mathbf{k}}^{\alpha\dagger} t_{\mathbf{k}}^\alpha + \Delta_{\mathbf{k}} (t_{\mathbf{k}}^{\alpha\dagger} t_{-\mathbf{k}}^{\alpha\dagger} + t_{\mathbf{k}}^\alpha t_{-\mathbf{k}}^\alpha)] + H_0$, where $H_0 = N(-3J_1 \tilde{s}^2/4 - \mu \tilde{s}^2 + \mu)$, $\Gamma_{\mathbf{k}} = J_1/4 - \mu - (J_2/2) \tilde{s}^2 \cos(k_x a) + J_3 \tilde{s}^2 \cos(k_y b)$, and $\Delta_{\mathbf{k}} = -(J_2/4) \tilde{s}^2 \cos(k_x a) + (J_3/2) \tilde{s}^2 \cos(k_y b)$, with $a = a_1 + a_2$.

By using the Bogliubov transformation, the Hamiltonian can be diagonalized,

$$H = \sum_{\mathbf{k}} \omega_{\mathbf{k}} \gamma_{\mathbf{k}}^{\alpha\dagger} \gamma_{\mathbf{k}}^\alpha + E. \quad (6)$$

The zero-temperature free energy is $E = N(-\frac{3J_1}{4} \tilde{s}^2 - \mu \tilde{s}^2 + \mu) + \frac{N}{(2\pi)^2} \int \int_{-\pi}^{\pi} dk_x dk_y \frac{3}{2} (\Omega_{\mathbf{k}} - \Lambda_{\mathbf{k}})$, where $\Omega_{\mathbf{k}} = (J_1/4 - \mu) \sqrt{1 - m \cos k_x + n \cos k_y}$, $\Lambda_{\mathbf{k}} = (J_1/4 - \mu) [1 - (m/2) \cos k_x + (n/2) \cos k_y]$, with $m = J_2 \tilde{s}^2 / (J_1/4 - \mu)$ and $n = 2J_3 \tilde{s}^2 / (J_1/4 - \mu)$. The energy spectrum can be expressed as

$$\omega_{\mathbf{k}} = (J_1/4 - \mu) \sqrt{1 - m \cos k_x a + n \cos k_y b}. \quad (7)$$

Minimizing the free energy with respect to \tilde{s} and μ gives $\partial E / \partial \tilde{s} = 0$ and $\partial E / \partial \mu = 0$, which determines the parameters \tilde{s} and μ self-consistently. The energy gap between the ground state and the lowest point of spectrum is

$$\Delta = (J_1/4 - \mu) \sqrt{1 - (m + n)}. \quad (8)$$

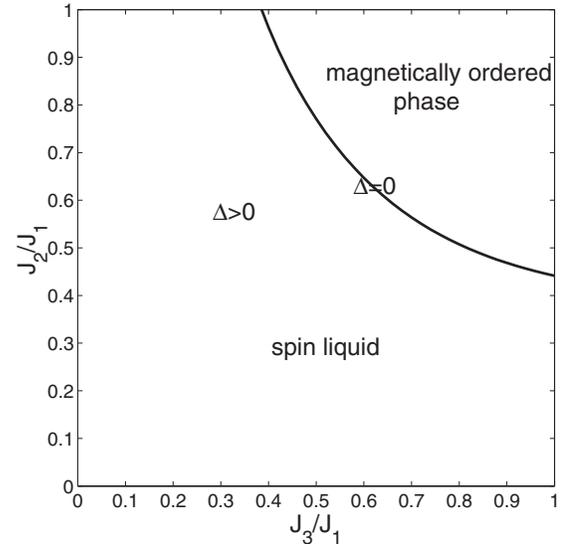


FIG. 3. Phase diagram controlled by the spin coupling strength. In the regime of the energy gap, $\Delta > 0$, the system is spin disordered or in a spin liquid phase, while a magnetically ordered phase appears at $\Delta = 0$. The phase boundary can be determined by the equation $(J_2 + 2J_3) \tilde{s}^2 = J_1/4 - \mu$, where the amplitude of condensate \tilde{s} and chemical potential μ can be given by the variational method. J_1, J_2 , and J_3 are functions of the lattice parameters, so the phase diagram is described by the amplitudes of standing waves in Fig. 4.

In the regime that $\Delta > 0$, most spins form spin singlets in pairs. In this regime the system is spin disordered or named as spin liquid [26]. The three triplet modes are degenerate and massive in the spin liquid phase, as displayed in Fig. 2. When the inter-dimer coupling increases, the gap between excited and ground states decreases and the spectrum is more dispersive. At a critical point, the energy gap vanishes at the point $\mathbf{k}_0 = (0, \pi/b)$ in the wave vector space. The t_α bosons condense and a magnetically ordered phase arises. The phase diagram of the spin liquid and magnetically ordered states is presented in Figs. 3 and 4. In Fig. 3, the controlling parameters are the ratio between the intradimer coupling and inter-dimer one. Due to the limitation of the bond operator method, the mean-field phase diagram needs to be improved when J_2/J_1 approaches 1 and J_3/J_1 approaches 0. In Fig. 4, we gave the phase diagram in terms of lattice parameters. Adjusting the intensity of lasers, it is possible to realize the transition between two phases in optical lattices.

In the magnetically disordered phase, the excitation spectra of triplet bosons are threefold degenerate. Considering spontaneous symmetry breaking, we assume that t_z bosons condense as the energy gap Δ approaches zero. The ground state of a rung in the ordered phase can be expressed as $|\tilde{\Phi}_0\rangle = \frac{1}{\sqrt{1+\lambda^2}} \prod_{i,j} (s_{i,j}^\dagger + \lambda e^{i\mathbf{k}_0 \cdot \mathbf{R}_{i,j}} t_{i,j}^\dagger) |0\rangle$, where λ is the condensation amplitude of triplet bosons and \mathbf{k}_0 represents the in-plane ordering wave vector (the gap begins to vanish in this point). The expected values of spin in this ground state are $\langle (S_{i,j}^l)_x \rangle = \langle (S_{i,j}^l)_y \rangle = \langle (S_{i,j}^l)_z \rangle = 0$ and $\langle (S_{i,j}^l)_z \rangle = -\langle (S_{i,j}^l)_z \rangle = [\lambda / (1 + \lambda^2)] \cos(\mathbf{k}_0 \cdot \mathbf{R}_{i,j})$. It is easy to see that all the dimers have a staggered magnetic structure and the magnitude of the spin expectation is related to the condensation of t_α bosons.

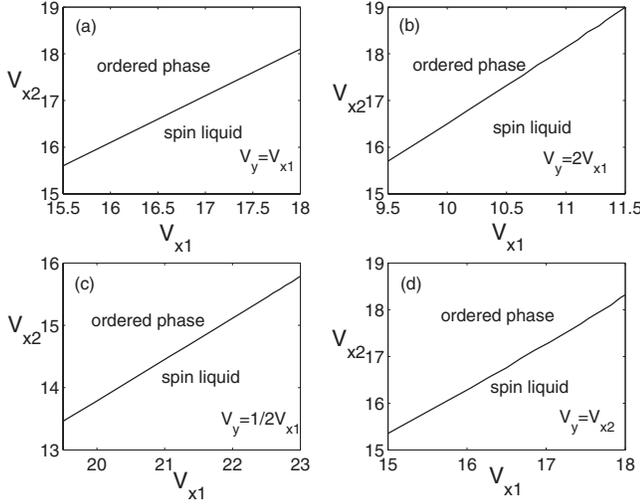


FIG. 4. Phase diagram in the V_{x1} - V_{x2} plane for different V_y . The wave vector satisfies $k_x = 2k_y$. V_{x1} , V_{x2} , and V_y are rescaled by the recoil energy E_r .

In order to describe the excitations of the ordered phase, we take the transformation $s'_{i,j} = (s_{i,j} + \lambda e^{i\mathbf{k}_0 \cdot \mathbf{R}_{i,j}} t'_{i,j}) / \sqrt{1 + \lambda^2}$, $t'_{i,j} = (-\lambda e^{i\mathbf{k}_0 \cdot \mathbf{R}_{i,j}} s_{i,j} + t_{i,j}) / \sqrt{1 + \lambda^2}$, and $t'_{i,j} = t_{i,j}$, $t'_{i,j} = t_{i,j}$. By performing this transformation and taking the Holstein-Primakoff expansion, $s'_{i,j} = s_{i,j} + (1 - 1/N) t'_{i,j} + \dots$ [21], the Hamiltonian includes zero-, linear-, and quadratic-order terms in the t' bosonic operators. The disappearance of the linear term gives the value of $\lambda = (J_2 + 2J_3 - J_1) / (J_2 + 2J_3 + J_1)$, which also minimizes the zero-order term. Finally, the Hamiltonian can be diagonalized as

$$H = \sum_{\mathbf{k}\alpha} \omega_{\mathbf{k}}^{\alpha} \gamma_{\mathbf{k}}^{\alpha\dagger} \gamma_{\mathbf{k}}^{\alpha} + \sum_{\mathbf{k}} \left(\omega_{\mathbf{k}}^x + \frac{1}{2} \omega_{\mathbf{k}}^z - A_{\mathbf{k}} - \frac{1}{2} B_{\mathbf{k}} \right) + E_0, \quad (9)$$

where $\omega_{\mathbf{k}}^x = \omega_{\mathbf{k}}^y = \sqrt{A_{\mathbf{k}}^2 - 4C_{\mathbf{k}}^2}$, $\omega_{\mathbf{k}}^z = \sqrt{B_{\mathbf{k}}^2 - 4D_{\mathbf{k}}^2}$, $E_0 = N(J_1/4)(\lambda^2 - 3)/(1 + \lambda^2) - \lambda^2(J_2 + 2J_3)/(1 + \lambda^2)^2$, and $A_{\mathbf{k}} = \frac{J_1}{(1 + \lambda^2)^2} + (J_2 + 2J_3) \frac{2\lambda^2}{(1 + \lambda^2)^2} + \frac{1 - \lambda^2}{1 + \lambda^2} M$, $B_{\mathbf{k}} = J_1 \frac{1 - \lambda^2}{1 + \lambda^2} + (J_2 + 2J_3) \frac{4\lambda^2}{(1 + \lambda^2)^2} + \frac{4\lambda^2}{(1 + \lambda^2)^2} M$, and $C_{\mathbf{k}} = \frac{M}{2}$, $D_{\mathbf{k}} = \frac{(1 - \lambda^2)^2 M}{(1 + \lambda^2)^2}$, with $M = -(J_2/2) \cos(k_x a) + J_3 \cos(k_y b)$. As a result, the energy spectrum in the magnetically ordered phase is obtained, as displayed in Fig. 5. In this phase, there are two degenerate spin-wave excitations restoring the breaking of rotational symmetry and one longitudinal mode which is gapped and corresponds to fluctuations in the amplitude of the moment.

IV. DETECTION OF THE PHASE TRANSITION

In the spin liquid and magnetically ordered phases, there are gapped excitations which are triplet magnons in spin liquid and longitudinal modes in the ordered phase. These gaps can be probed by magnetic resonance [8]. When $\mu_B B \ll J_1$ (μ_B is the magnetic moment of the alkali-metal atom and B is the intensity of the oscillating magnetic field), the frequency of oscillating magnetic field has a resonant peak at $\omega = E_{\text{gap}}/\hbar$.

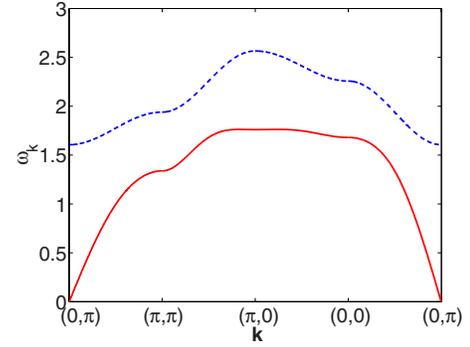


FIG. 5. (Color online) Energy spectrum in the magnetically ordered phase. The upper dashed line is the longitudinal mode, and the lower solid one is the doubly degenerate transverse mode. The coupling strength $J_2/J_1 = J_3/J_1 = 0.7$. $\lambda = 0.6$. The corresponding lattice parameters are $V_{x1} = 27E_r$, $V_{x2} = 16E_r$, and $V_y = 10E_r$. $\omega_{\mathbf{k}}$ is measured in unit of J_1 . k_x and k_y are measured in units of $1/a$ and $1/b$, respectively.

The excited spectrum is also a sign to distinguish the two phases. Two-photon Bragg scattering provides an effective method to measure the spectrum [3,27]. In such experiments two laser beams with different wave vectors and frequencies illuminate the atom cloud. The frequency difference is much smaller than the detuning of the two lasers from atomic resonance. The atoms absorb a photon from one beam and emit another photon into the other. By changing the angle between the two laser beams, we can tune the momentum transfer (momentum difference between the two beams). If the momentum and frequency differences match those of the dispersion relation of the spectrum, a resonant absorption of the probe light happens. Other methods for measuring spin correlation functions were proposed to distinguish the different magnetic phases [8].

V. CONCLUSION

In summary, 2D coupled spin ladders can be simulated by an optical superlattice. A phase transition from the spin liquid to magnetically ordered phase is possibly realized through adjusting the lattice parameters controlled by laser beams. Our results show that strongly correlated cold atoms in optical lattices provide a route to observe this quantum phase transition. Manipulating the amplitudes and wave vectors of laser beams, the coupling between spins can be adjusted in a wide range. Recent developments of controlling the cold atoms in optical lattices allow for the experimental investigation of our prediction in the future.

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