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Spin dynamics of a J_1 - J_2 -K model for the paramagnetic phase of iron pnictides

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We study the finite-temperature spin dynamics of the paramagnetic phase of iron pnictides within an antiferromagnetic J_1 - J_2 Heisenberg model on a square lattice with a biquadratic coupling $-K(\mathbf{S}_i \cdot \mathbf{S}_j)^2$ between the nearest-neighbor spins. Our focus is on the paramagnetic phase in the parameter regime of this J_1 - J_2 -K model where the ground state is a $(\pi, 0)$ collinear antiferromagnet. We treat the biquadratic interaction via a Hubbard-Stratonovich decomposition and study the resulting effective quadratic-coupling model using both modified spin wave and Schwinger boson mean-field theories; the results for the spin dynamics derived from the two methods are very similar. We show that the spectral weight of dynamical structure factor $S(\mathbf{q},\omega)$ is peaked at ellipses in the momentum space at low excitation energies. With increasing energy, the elliptic features expand towards the zone boundary and gradually split into two parts, forming a pattern around (π, π) . Finally, the spectral weight is anisotropic, being larger along the major axis of the ellipse than along its minor axis. These characteristics of the dynamical structure factor are consistent with the recent measurements of the inelastic neutron scattering spectra of BaFe₂As₂ and SrFe₂As₂.

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I. INTRODUCTION

The emergence of superconductivity in iron pnictides^{1,2} near an antiferromagnetically ordered state³ in the phase diagram suggests strong interplay between superconductivity and magnetism in these materials. Elucidating the magnetic excitations is therefore important for understanding not only the overall microscopic physics of these systems but also their superconductivity. In the parent compounds, the observed (π ,0) antiferromagnetic order arises either within a weakcoupling approach invoking a Fermi surface nesting^{4–6} or from a strong-coupling approach whose starting point is a local moment J_1 - J_2 -K model.^{7–14}

The strong-coupling approach is based on the proximity of the metallic ground state of the parent pnictides to a Mott localization transition, which gives rise to quasilocal magnetic moments.^{7,12,15,16} This incipient Mott picture corresponds to a ratio of U (a measure of the Coulomb repulsions and Hund's couplings among the Fe 3d electrons) to t (the characteristic bandwidth of the Fe 3d electrons); this is not too far below the Mott threshold U_c/t , which is usually of order unity. This is supported by many experimental observations. For instance, the room-temperature electrical resistivity of parent iron pnictides is so high (even when the residual resistivity is relatively low, signaling the smallness of elastic scattering) that the extracted mean-free path of quasiparticles would be comparable to the Fermi wavelength; this is typical of bad metals near a Mott trasition. Similarly, the Drude weight in optical conductivity^{17,18} is strongly suppressed from its noninteracting counterpart, providing a direct measure of the proximity to the Mott transition. This is further corroborated by the temperature-induced spectral weight transfer,¹⁸⁻²⁰ which is also characteristic of metals near a Mott transition. In the spin sector, zone boundary spin waves have been observed by inelastic neutron scattering (INS) measurements in the magnetically ordered state of several 122 iron pnictide compounds.²¹ Both the large spectral weight and the relatively small spin damping suggest quasilocalized moments, which are expected near the Mott transition, where the spin excitations arise out of incoherent electronic excitations. Additional evidence for the incipient Mott transition picture has come from the observation of a Mott insulating phase in iron oxychalcogenides.²² This material contains an expanded Fe square lattice compared to the iron pnictides, which reduces *t*, thereby enhancing U/tbeyond U_c/t .²² Likewise, the Mott insulating behavior of the alkaline iron selenides²³ can also be interpreted as the result of a reduced effective *t* and, correspondingly, an enhanced U/tbeyond U_c/t .^{24,25}

In the vicinity of U_c/t , where correlations are strong, it is natural that the spin Hamiltonian contains not only two-spin interactions, such as J_1 and J_2 Heisenberg exchange between nearest- and next-nearest-neighbor spins on a square lattice, but also interactions involving a higher number of spins. These naturally include, for instance, the ring-exchange coupling involving four spins on a plaquette and the biquadratic coupling of the form $-K(\mathbf{S}_i \cdot \mathbf{S}_j)^2$ in systems of spin size $S \ge 1.^{26}$ The subject of the present study is to show how such non-Heisenberg interactions, particularly the biquadratic interaction, influence the spin dynamics in the paramagnetic phase.

Spin dynamics in the parent iron pnictides have been most extensively studied in the low-temperature state $(T < T_N)$ with both antiferromagnetic order and orthorhombic structural distortion. Here, the INS experiments up to high energies (of the order of 200 meV) show that the spin-wave excitations in these compounds are highly anisotropic, with a dispersion which can be understood in terms of an anisotropic J_{1x} - J_{1y} - J_2 model with $J_{1x} \neq J_{1y}$.^{21,27,28} The anisotropy in the nearest-neighbor coupling is compatible with an orthorhombic structure, and its degree could reflect an orbital ordering.^{29–32} Detailed theoretical studies of the magnetic excitations in the ordered phase have been carried out in such a J_{1x} - J_{1y} - J_2 model,³³ and in a J_1 - J_2 -K model.^{34,35} It should also be noted that terms such as the biquadratic coupling could be inferred from the sublattice angle dependence of the ground-state energy in LSDA calculations³⁶ and can also appear naturally as a result of the orbital ordering between Fe d_{xz} and d_{yz} orbitals.²⁹

Our focus is instead on the spin dynamics in the *paramagnetic phase* of the parent iron pnictides, which has only recently been studied experimentally. The initial work by Diallo *et al.*³⁷ measured the spin dynamics of CaFe₂As₂ at relatively low energies, below 70 meV. Theoretically, four of us³⁸ studied the spin dynamics in the paramagnetic phase of the J_1 - J_2 model (with or without an additional fermion damping). We showed that the experimentally observed elliptical features of the spin spectral weight in momentum space are well described by this model and we determined the change to the elliptical features at high energies.

More recently, Harriger et al.39 reported measurements of the spin dynamics in the paramagnetic phase up to high energies (above 200 meV) in BaFe2As2. The INS measurements confirmed the quasi-two-dimensional (2D) spin dynamics found at low energies³⁷ and characterized the evolution of the low-energy elliptic features as they expand towards the zone boundary as the energy is raised and determined the high-energy dispersion, which appears to require a $J_{1x} \neq J_{1y}$ description even though the paramagnetic phase has a tetragonal structure. Similar data have also been reported by Ewings et al. in SrFe₂As₂.⁴⁰ Theoretically, Park *et al.*⁴¹ analyzed the spin dynamics in the paramagnetic state within a dynamical mean-field theory (DMFT) for interactions $U/t \lesssim U_c/t$, demonstrating that the DMFT approach captures key features of the neutron scattering results, including the ellipticity of the map of the structure-factor peak in the Brillouin zone.

In this paper, we study the spin dynamics of the J_1 - J_2 -K model in the tetragonal paramagnetic phase using both modified spin-wave (MSW) and Schwinger boson mean-field (SBMF) theories. The results from the two methods are in very good quantitative agreement with each other. We show that, for a moderate biquadratic coupling K, the dynamical structure factor $S(\mathbf{q},\omega)$ has not only elliptic features near $(\pi,0)$, which expand with increasing energy and split into peaks surrounding (π,π) , but also an anisotropic distribution of the spectral weight that is larger along the major axis of each ellipse than along its minor axis. These properties agree well with the INS experiments^{39,40}

The remainder of the paper is organized as follows. In Sec. II we introduce the J_1 - J_2 -K model and describe the MSW and SBMF theories used in this paper. In Sec. III we show how the biquadratic coupling K influences the mean-field phase diagram and magnetic excitation spectrum. In Sec. IV we calculate the dynamical structure factor $S(\mathbf{q},\omega)$. We also show that the spectral weight exhibits anisotropic features, discuss the evolution of the anisotropic features with increasing excitation energy, and explain how these properties arise from our theory. In Sec. V we first discuss some possible generalizations of the J_1 - J_2 -K model study in this paper. In the same section, we then consider the effect of itinerant electrons and compare our study with other theoretical approaches to the spin dynamics. Section VI is devoted to a comparison with INS experiments on the paramagnetic phases of the parent 122 iron pnictides, and Sec. VII contains a few concluding remarks. In the three Appendixes, we expound on the Ising transition at

low J_1/J_2 ratios and discuss the effects of both ring-exchange interactions and interlayer exchange couplings.

II. MODEL AND METHODS

The J_1 - J_2 -K model is defined on a 2D square lattice with the following Hamiltonian:

$$H = J_1 \sum_{i,\delta} \mathbf{S}_i \cdot \mathbf{S}_{i+\delta} + J_2 \sum_{i,\delta'} \mathbf{S}_i \cdot \mathbf{S}_{i+\delta'} - K \sum_{i,\delta} (\mathbf{S}_i \cdot \mathbf{S}_{i+\delta})^2,$$
(1)

where J_1 and J_2 , respectively, denote the antiferromagnetic exchange couplings between spins located in the nearestneighbor ($\delta = \hat{x}, \hat{y}$) and next-nearest-neighbor ($\delta' = \hat{x} \pm \hat{y}$) sites. *K* is the coupling for the biquadratic interaction between the nearest-neighbor spin pairs.

To fully explain the experimentally observed $(\pi,0,\pi)$ antiferromagnetic order, an exchange coupling along the third dimension, J_z , should also be included. However, we find that the model defined in Eq. (1) already allows us to understand the experimentally observed quasi-2D spin dynamics. Hence, we concentrate on this 2D model in the text and discuss the influence of the interlayer coupling J_z on the spin dynamics in Appendix C.

The Hamiltonian of Eq. (1) is studied using both MSW^{42,43} and SBMF⁴⁴ methods. Here, we focus on the parameter regime where the ground state has a collinear (π ,0) antiferromagnetic order and decompose the biquadratic interaction term of the Hamiltonian using two Hubbard-Stratonovich fields $\Gamma_{i,\hat{x}(\hat{y})}$. The effective Hamiltonian reads

$$H = J_1 \sum_{i,\delta} \mathbf{S}_i \cdot \mathbf{S}_{i+\delta} + J_2 \sum_{i,\delta'} \mathbf{S}_i \cdot \mathbf{S}_{i+\delta'} - 2K \sum_{i,\delta} \Gamma_{i,\delta} \mathbf{S}_i \cdot \mathbf{S}_{i+\delta} + K \sum_{i,\delta} \Gamma_{i,\delta}^2.$$
(2)

At the mean-field level, the Hubbard-Stratonovich fields are treated as static quantities and can be expressed using equaltime spin correlators as $\Gamma_{i,\delta} = \langle \mathbf{S}_i \cdot \mathbf{S}_{i+\delta} \rangle$. The static approximation is made in accordance with the level of approximation inherent to the MSW and SBMF methods, which incorporate static self-energies for the respective boson fields. As shown below, our approach has two important features: (i) it is capable of studying the Ising correlations at nonzero temperatures; and (ii) the MSW and SBMF approaches yield consistent results.

A. The modified spin-wave theory

The MSW theory^{42,43} has been adapted to the J_1 - J_2 model by four of us.³⁸ In this approach, a local spin quantization axis is defined at each site along the classical ordering direction Ω_i^{cl} . The Hamiltonian in Eq. (2) is then expressed in terms of Dyson-Maleev (DM) bosons via a local DM transformation: $\mathbf{S}_i \cdot \Omega_i^{cl} = S - a_i^{\dagger} a_i$, $\mathbf{S}_i^+ = \sqrt{2S}(1 - a_i^{\dagger} a_i/2S)a_i$, and $\mathbf{S}_i^- = \sqrt{2S}a_i^{\dagger}$. Minimizing the free energy under the constraint of zero sublattice magnetization $\langle S - a_i^{\dagger} a_i \rangle = 0$ by introducing a Lagrange multiplier μ , and with respect to $\Gamma_{\delta} (= \Gamma_{i,\delta}$, by assuming translational symmetry), we obtain

$$\Gamma_{x} = \cos^{2} \frac{\phi}{2} f_{x}^{2} - \sin^{2} \frac{\phi}{2} g_{x}^{2},$$
(3)

$$\Gamma_{y} = \sin^{2} \frac{\phi}{2} f_{y}^{2} - \cos^{2} \frac{\phi}{2} g_{y}^{2},$$

where $\phi = \arccos(\Omega_i^{cl} \cdot \Omega_{i+\hat{x}}^{cl})$. $f_{\delta} = \langle a_i^{\dagger} a_{i+\delta} \rangle$ and $g_{\delta} = \langle a_i a_{i+\delta} \rangle$ are the ferromagnetic and antiferromagnetic bond operators, respectively. Minimizing the free energy with respect to ϕ gives $\sin \phi = 0$ for nonzero f_{δ} and g_{δ} , or ϕ can be arbitrary if $f_{\delta} = g_{\delta} = 0$. This defines two phases separated by a mean-field temperature scale³⁸ $T_{\sigma 0}$: at $T > T_{\sigma 0}$, ϕ is arbitrary, and the system has C_{4v} lattice rotational symmetry; while at $T < T_{\sigma 0}$, the C_{4v} symmetry is broken and the system is Ising ordered, corresponding to either $\phi = 0$ or $\phi = \pi$. In MSW theory, the Ising order parameter can be defined as $\sigma = 2(\cos^2 \frac{\phi}{2}(f_x^2 + g_y^2) - \sin^2 \frac{\phi}{2}(f_y^2 + g_x^2))$. From Eq. (3), if we define $\Gamma_{\pm} = (\Gamma_x \pm \Gamma_y)/2$ as the symmetric and antisymmetric Hubbard-Stratonovich fields, we find that $\Gamma_{-} = \sigma/4$.

Minimizing the free energy with respect to other variational parameters, we obtain a set of self-consistent equations:

$$f_{\delta} = m_0 + \frac{1}{N} \sum_{\mathbf{k}}' \frac{B_{\mathbf{k}}}{\varepsilon_{\mathbf{k}}} \left(n_{\mathbf{k}} + \frac{1}{2} \right) \cos(\mathbf{k} \cdot \boldsymbol{\delta}), \qquad (4)$$

$$g_{\delta} = m_0 + \frac{1}{\mathcal{N}} \sum_{\mathbf{k}}' \frac{A_{\mathbf{k}}}{\varepsilon_{\mathbf{k}}} \left(n_{\mathbf{k}} + \frac{1}{2} \right) \cos(\mathbf{k} \cdot \boldsymbol{\delta}'), \qquad (5)$$

$$S + \frac{1}{2} = m_0 + \frac{1}{\mathcal{N}} \sum_{\mathbf{k}}' \frac{B_{\mathbf{k}}}{\varepsilon_{\mathbf{k}}} \left(n_{\mathbf{k}} + \frac{1}{2} \right), \tag{6}$$

where \mathcal{N} is the total number of lattice sites, $\delta = \hat{x}, \hat{y}$, and $\delta' = \hat{x}, \hat{y}, \hat{x} \pm \hat{y}$. In Eqs. (4)–(6),

$$A_{\mathbf{k}} = 2\sin^2 \frac{\phi}{2} \tilde{J}_{1x} g_x \cos k_x + 2\cos^2 \frac{\phi}{2} \tilde{J}_{1y} g_y \cos k_y + 4J_2 g_{x+y} \cos k_x \cos k_y,$$
(7)

$$B_{\mathbf{k}} = 4J_2g_{x+y} - \mu + 2\sin^2\frac{\phi}{2}(\tilde{J}_{1x}g_x - \tilde{J}_{1y}f_y(1 - \cos k_y))$$

$$+ 2\cos^2\frac{\phi}{2}(\tilde{J}_{1y}g_y - \tilde{J}_{1x}f_x(1 - \cos k_x)), \tag{8}$$

and the Bogoliubov angle $\theta_{\mathbf{k}}$ is defined via $\tanh 2\theta_{\mathbf{k}} = A_{\mathbf{k}}/B_{\mathbf{k}}$. The boson dispersion $\varepsilon_{\mathbf{k}} = \sqrt{B_{\mathbf{k}}^2 - A_{\mathbf{k}}^2}$ and the boson number $n_{\mathbf{k}} = [\exp(\varepsilon_{\mathbf{k}}/k_BT) - 1]^{-1}$. At T = 0, the spectrum of the DM bosons becomes gapless at wave vector \mathbf{Q} and $\mathbf{0}$. This corresponds to a long-range antiferromagnetic order at $\mathbf{Q} \neq 0$ with a nonzero spontaneous magnetization m_0 . In this case, the summation $\sum_{\mathbf{k}}'$ runs over all \mathbf{k} values that make $\varepsilon_{\mathbf{k}} > 0$, and the contribution from the $\varepsilon_{\mathbf{k}} = 0$ terms is taken into account separately by m_0 . For T > 0, $m_0 = 0$, and the system is paramagnetic. Here the summation is performed in the full momentum space. In the presence of a small third-dimension coupling J_z , there will be a nonzero mean-field Néel temperature, T_{N0} ; this is discussed in Appendix C.

Note that these self-consistent equations are exactly the same as those for the isotropic $J_1 - J_2$ model.³⁸ But the definitions of A_k and B_k are different. In Eqs. (7) and (8) above, we defined the effective exchange couplings J_{1x} (J_{1y})

along the x(y) direction as

$$\tilde{J}_{1x(y)} = J_1 - 2K\Gamma_{x(y)},$$
(9)

expressed in terms of the Hubbard-Stratonovich fields $\Gamma_{x(y)}$ of spin-spin correlators in Eq. (3). Although in the J_1 - J_2 -K model the bare nearest-neighbor exchange coupling J_1 is still isotropic, a nonzero biquadratic coupling K leads to an anisotropic *effective coupling* $\tilde{J}_{1x} \neq \tilde{J}_{1y}$ in the Isingordered phase where $\Gamma_x \neq \Gamma_y$; i.e., the nearest-neighbor spin correlators along x and y are unequal, similarly to the situation found originally⁴⁹ for the J_1 - J_2 model.

B. The Schwinger boson theory

In the Schwinger boson representation,⁴⁴ the SU(2) spin operators are rewritten in terms of two Schwinger bosons via the transformation $S_i^z = \frac{1}{2}(a_i^{\dagger}a_i - b_i^{\dagger}b_i)$, $S_i^+ = a_i^{\dagger}b_i$, and $S_i^- = b_i^{\dagger}a_i$. To limit the boson Hilbert space to the physical sector, a constraint $a_i^{\dagger}a_i + b_i^{\dagger}b_i = 2S$ is imposed on each site. This can be generalized to the case of either SU(*N*)⁴⁵ or SP(*N*)^{46,47} spins; in either case there will be *N* boson degrees of freedom at each site. For the experimentally observed $\mathbf{Q} = (\pi, 0)$ or $(0, \pi)$ antiferromagnetic collinear phase in the 122 parent compounds, the (ab)-plane spin-spin correlations are expressed as

$$\mathbf{S}_{i} \cdot \mathbf{S}_{j} = -(1 - \Theta(i, j))[2\hat{g}_{ij}^{\dagger}\hat{g}_{ij} - S^{2}] + \Theta(i, j)[2\hat{f}_{ij}^{\dagger}\hat{f}_{ij} - S(S+1)], \qquad (10)$$

where $\hat{f}_{\delta} \equiv f_{i,i+\delta} = \frac{1}{2}(a_i^{\dagger}a_{i+\delta} + b_i^{\dagger}b_{i+\delta})$ and $\hat{g}_{\delta} \equiv g_{i,i+\delta} = \frac{1}{2}(a_ib_{i+\delta} - b_ia_{i+\delta})$ are, respectively, the ferromagnetic and antiferromagnetic bond operators. The function $\Theta(i, j) = 1$ if *i* and *j* are on the same stripe sublattice, and $\Theta(i, j) = 0$ if *i* and *j* are on different stripe sublattices. The Hubbard-Stratonovich field is then $\Gamma_{\delta} = |\hat{f}_{\delta}|^2 - |\hat{g}_{\delta}|^2$, and in the case of $(\pi, 0)$ ordering we find

$$\Gamma_x = -g_x^2, \quad \Gamma_y = f_y^2. \tag{11}$$

Comparing this to Eq. (3), we see that the spin correlators coincide with those in the MSW theory if one sets $\phi = \pi$. Similarly, the case of $(0,\pi)$ ordering corresponds to $\phi = 0$ in Eq. (3). In both cases, Γ_x and Γ_y have opposite sign, leading to the anisotropy in the effective spin-spin exchange couplings $\tilde{J}_{1x} \neq \tilde{J}_{1y}$, from Eq. (9).

By introducing Fourier transformation,⁴⁸

$$a_i = \frac{1}{\sqrt{N}} \sum_{\mathbf{k}} a_{\mathbf{k}} e^{i(\mathbf{k} - \frac{\mathbf{Q}}{2}) \cdot \mathbf{r}_i}, \qquad (12)$$

$$b_i = \frac{1}{\sqrt{N}} \sum_{\mathbf{k}} b_{\mathbf{k}} e^{i(\mathbf{k} + \frac{\mathbf{Q}}{2}) \cdot \mathbf{r}_i},\tag{13}$$

and making a Bogoliubov transformation to a new quasiparticle creation/annihilation operators $\alpha_k = \cosh \theta_k a_k + i \sinh \theta_k b_{-k}^{\dagger}$, one arrives at the mean-field free energy density, which can be generalized to the Sp(N) form⁴⁷

$$F_{\rm MF} = \frac{NT}{N} \sum_{\mathbf{k}} \ln\left[2\sinh\left(\frac{\omega_{\mathbf{k}}}{2T}\right)\right] + N\lambda\left(S + \frac{1}{2}\right) \\ -\frac{Nz}{8} \sum_{\delta} J_{\delta}(|f_{\delta}|^2 - |g_{\delta}|^2), \tag{14}$$

where z is the coordination number, and λ is the Lagrange multiplier associated with the imposed constraint that, on average, the number of bosons per site $\sum_{\sigma=1}^{N} n_{i\sigma} = NS$. Here $\omega_{\mathbf{k}} = \sqrt{(B_{\mathbf{k}}-\lambda)^2 - A_{\mathbf{k}}^2}$ is the dispersion of the Bogoliubov quasiparticles, expressed⁴⁸ in terms of the variables

$$A_{\mathbf{k}} = i \sum_{\delta} J_{\delta} g_{\delta} e^{-i\mathbf{k}\cdot\delta}, \quad B_{\mathbf{k}} = \sum_{\delta} J_{\delta} f_{\delta} e^{-i\mathbf{k}\cdot\delta}; \quad (15)$$

the Bogoliubov angle $\tanh 2\theta_{\mathbf{k}} = A_{\mathbf{k}}/(\lambda - B_{\mathbf{k}})$. The dispersion relation $\omega_{\mathbf{k}}$ explicitly depends on the ordering wave vector \mathbf{Q} and has minima around $\mathbf{k} = \pm \mathbf{Q}/2$. In the regime where $J_2 > J_1/2$, the minimization of the free energy results in $\mathbf{Q} = (\pi, 0)$ or $(0, \pi)$. For example, for $\mathbf{Q} = (\pi, 0)$, the expressions for $A_{\mathbf{k}}$ and $B_{\mathbf{k}}$ become

$$A_{\mathbf{k}} = 2\tilde{J}_{1x}g_x \sin k_x + 4J_2g_{x+y} \sin k_x \cos k_y, \qquad (16)$$

$$B_{\mathbf{k}} = 2\tilde{J}_{1y}f_y\cos k_y. \tag{17}$$

In the large-*N* limit of the Sp(*N*) spin, the mean-field free energy, Eq. (14), becomes exact.^{44,47} The observable magnetic excitation spectrum is obtained from $\omega_{\mathbf{k}}$ by a **Q**/2 shift: $\varepsilon_{\mathbf{k}} = \omega_{\mathbf{k}-\mathbf{Q}/2}$. At T = 0, the magnetic order results in the gapless Goldstone modes at $\mathbf{k} = \mathbf{0}$ and **Q**, as expected. The SBMF theory is known to reproduce well the spectrum of spin waves in both ferro- and antiferromagnets.^{44,48}

Below, we focus on the paramagnetic phase at T > 0, with short-range $\mathbf{Q} = (\pi, 0)$ antiferromagnetic correlations [the case $\mathbf{Q} = (0, \pi)$ is obtained by a C_4 lattice rotation]. We obtain the following self-consistent equations from the saddle-point minimization of the free energy, Eq. (14):

$$f_{\delta} = \frac{1}{\mathcal{N}} \sum_{\mathbf{k}} \frac{B_{\mathbf{k}} - \lambda}{\omega_{\mathbf{k}}} \left(n_{\mathbf{k}} + \frac{1}{2} \right) \cos\left(\mathbf{k} \cdot \boldsymbol{\delta}\right), \qquad (18)$$

$$g_{\delta'} = \frac{1}{\mathcal{N}} \sum_{\mathbf{k}} \frac{A_{\mathbf{k}}}{\omega_{\mathbf{k}}} \left(n_{\mathbf{k}} + \frac{1}{2} \right) \sin(\mathbf{k} \cdot \boldsymbol{\delta}'), \tag{19}$$

$$S + \frac{1}{2} = \frac{1}{\mathcal{N}} \sum_{\mathbf{k}} \frac{B_{\mathbf{k}} - \lambda}{\omega_{\mathbf{k}}} \left(n_{\mathbf{k}} + \frac{1}{2} \right), \tag{20}$$

where $\delta = \hat{y}$, and $\delta' = \hat{x}, \hat{x} \pm \hat{y}$. Under the transformation $B_{\mathbf{k}} - \lambda \rightarrow B_{\mathbf{k}}$ and $\mathbf{k} \rightarrow \mathbf{k} - \mathbf{Q}/2$, Eqs. (16)–(20) in the SBMF theory and Eqs. (4)–(8) in the MSW mean-field theory have exactly the same form in the short-range (π ,0) correlated paramagnetic phase. Therefore, the two methods yield exactly the same mean-field phase diagram and boson dispersion, as corroborated by explicit numerical comparison. We further verified that these two theories give similar results for the spin dynamics of the J_1 - J_2 -K model.



FIG. 1. (Color online) Mean-field phase diagram in the MSW theory for S = 1 and $J_1/J_2 = 1$. Phases I, II, and III, respectively, denote the $(\pi, 0)/(0, \pi)$ long-range antiferromagnetically ordered state (at T = 0), the Ising-ordered paramagnetic state, and the isotropic paramagnetic state. The solid (red) curve refers to the mean-field temperature scale $T_{\sigma 0} = T_0$. In the shaded region the effective exchange coupling $\tilde{J}_{1y} < 0$.

III. MEAN-FIELD PHASE DIAGRAM AND EXCITATION SPECTRUM

Since INS measurements suggest $J_1 \sim J_2$ for several 122 compounds, 21,37,39 our discussion of the J_1 - J_2 -K model is focused on this parameter regime. Figure 1 shows the meanfield phase diagram of the 2D J_1 - J_2 -K model using the MSW method for S = 1 and $J_1/J_2 = 1$. We identify three phases. Phase I corresponds to the $(\pi, 0)/(0, \pi)$ antiferromagnetically long-range-ordered phase; it exists only at T = 0 in the 2D model. Phase II and phase III are both paramagnetic. They are separated by a mean-field Ising transition temperature $T_{\sigma 0}$. We find that for $J_1/J_2 = 1$, this transition is first-order, as shown in Fig. 2. But it can be either first-order or second-order for $J_1/J_2 \lesssim 0.9$, as discussed in more detail in Appendix A. In the low-temperature phase II, either $f_x \neq f_y$ or $g_x \neq g_y$ (see Fig. 2), corresponding to an Ising-ordered phase with either $(\pi,0)$ or $(0,\pi)$ short-range antiferromagnetic correlations. This Ising-ordered phase already exists in the isotropic J_1 - J_2 model.^{38,47,49} But here, we find that a nonzero K enhances $T_{\sigma 0}$, and K drives the effective nearest-neighbor exchange couplings to be anisotropic. As shown in Fig. 2, in the $(\pi, 0)$ Ising-ordered phase (corresponding to $\phi = \pi$), the effective coupling \tilde{J}_{1y} can even be ferromagnetic. This is important for understanding the experimentally observed anisotropic magnetic excitations at high energies in Ca-122²¹ and Ba-122.³⁹ Phase III at $T > T_{\sigma 0}$ is the Ising-disordered paramagnetic phase. In this phase the effective nearest-neighbor exchange couplings are isotropic because the nearest-neighbor bond correlators are 0. But the next-nearest-neighbor bond correlations may still be finite in this phase. One may define another temperature scale T_0 , above which the next-nearestneighbor bond correlations vanish and the system is decoupled into isolated local moments. Note that T_0 does not refer to a phase transition, and the discontinuity of the bond correlations at T_0 is an artifact of the mean-field theory.⁴² In general, T_0 and $T_{\sigma 0}$ are two different temperature scales satisfying $T_0 \ge T_{\sigma 0}$.^{38,47} But for $J_1/J_2 \gtrsim 0.9$, $T_0 = T_{\sigma 0}$ for any K/J_2



FIG. 2. (Color online) Temperature evolution of the mean-field parameters in the MSW theory for S = 1, $J_1/J_2 = 1$, and $K/J_2 = 0.8$.

ratio, as shown in Fig. 2. The phase diagram obtained in the SBMF theory is identical to the one shown in Fig. 1.

The finite coupling *K* not only changes the phase boundary of the mean-field phase diagram, but also can dramatically influence the boson excitation spectrum. In Fig. 3(a), we show the dispersions of the DM and Schwinger bosons along two high-symmetry directions in momentum space for various *K* values in phase II with $\phi = \pi$ using the same parameters as in Fig. 1. We see that the dispersion in Schwinger boson theory matches the one in the MSW theory exactly.

The dispersion shows a gap at $(\pi, 0)$ [and also at (0, 0)], with the size

$$\Delta_1 = \sqrt{-\mu(8J_2g_{x+y} + 4\tilde{J}_{1x}g_x - \mu)}.$$
 (21)

At low temperatures the gap is small since $\mu \to 0$ as $T \ll T_{\sigma 0}$. In this limit, the excitation near $(\pi, 0)$ can be approximated by $\varepsilon_{\mathbf{k}} = \sqrt{v_{1x}^2 (\pi - k_x)^2 + v_{1y}^2 k_y^2 + \Delta_1^2}$, where the velocities are, respectively,

$$v_{1x} = 4J_2g_{x+y} + 2\tilde{J}_{1x}g_x, \qquad (22)$$

$$v_{1y} = \sqrt{(4J_2g_{x+y} + 2\tilde{J}_{1x}g_x)(4J_2g_{x+y} - 2\tilde{J}_{1y}f_y) + 2\tilde{J}_{1y}f_y\mu}.$$
(23)

The excitation develops to the gapless Goldstone mode at T = 0. At (π,π) [and also at $(0,\pi)$] the dispersion has a different gap:

$$\Delta_2 = \sqrt{(8J_2g_{x+y} - 4\tilde{J}_{1y}f_y - \mu)(4\tilde{J}_{1x}g_x - 4\tilde{J}_{1y}f_y - \mu)}.$$
(24)



FIG. 3. (Color online) (a) MSW dispersion along high-symmetry directions in the paramagnetic Brillouin zone for the 2D J_1 - J_2 -K model at S = 1, $J_1/J_2 = 1.0$, $T/J_2 = 1.0$, and various K values. For comparison, the dashed (red) curve shows the dispersion in the SBMF theory for the same parameters and $K/J_2 = 0.8$. The gaps at (0,0) and $(\pi, 0)$ are too small to be seen here. (b) Symbols show the dispersion from INS data at T = 150 K in BaFe₂As₂, taken from Ref. 39. The data can be fit by any of the theoretical dispersion curves [as in (a)] that lie within the shaded region.

The features that $v_{1x} \neq v_{1y}$ and $\Delta_1 \neq \Delta_2$ already exist in the isotropic J_1 - J_2 model. In the J_1 - J_2 -K model, Δ_1 at $(\pi, 0)$ is only weakly affected by K because it is dominated by μ . But Δ_2 at (π, π) is strongly influenced. It increases with K. For sufficiently large K, approximately where \tilde{J}_{1y} changes sign to be ferromagnetic (the shaded region in Fig. 1), the dispersion at (π, π) turns from a local minimum to a maximum, as shown in Fig. 3(a). Similar behavior in the spin-wave dispersion of the J_1 - J_2 -K model has also been discussed in Ref. 34 in the antiferromagnetic phase.

IV. DYNAMICAL STRUCTURE FACTOR

In order to investigate the magnetic excitations, which are directly accessible by INS measurements, we have calculated the magnetic structure factor $S(\mathbf{q}, \omega)$. Our main interest is to understand the experimentally observed anisotropic feature of the magnetic excitations in the paramagnetic phase above the Néel temperature. As discussed in Sec. III, in this temperature regime, the most relevant factor for the in-plane anisotropy is the Ising order. Therefore, we concentrate our discussion on the magnetic structure factor in phase II of the 2D J_1 - J_2 -K model. In this phase $S(\mathbf{q}, \omega)$ has the same form in both the

MSW and the Schwinger boson theories,

$$S(\mathbf{q},\omega) = 2\pi \frac{C}{N} \sum_{\mathbf{k}}' \sum_{s,s'=\pm 1} [\cosh(2\theta_{\mathbf{k}+\mathbf{q}} - 2\theta_{\mathbf{k}}) - ss'] \\ \times \delta(\omega - s\varepsilon_{\mathbf{k}+\mathbf{q}} - s'\varepsilon_{\mathbf{k}}) n_{\mathbf{k}+\mathbf{q}}^{s} n_{\mathbf{k}}^{s'}, \qquad (25)$$

where \sum' refers to the summation over the magnetic Brillouin zone corresponding to the $(\pi, 0)$ order, which is enclosed by $-\pi/2 \leq k_x \leq \pi/2$ and $-\pi \leq k_y \leq \pi$. $n_k^+ = n_k + 1$ and $n_k^- = n_k$. C = 1 in the MSW theory, and C = 3/2 in the Schwinger boson theory with SU(2) symmetry.⁵⁰ We see from Eq. (25) that the contribution to $S(\mathbf{q}, \omega)$ comes from two-boson processes. Hence, in general cases the peak of $S(\mathbf{q},\omega)$ does not follow the boson dispersion. But at low temperatures, the largest contribution to $S(\mathbf{q},\omega)$ in the summation over **k** comes from the term at $\mathbf{k} = (0,0)$ since the small gap Δ_1 at this point results in a large boson number n_k . To satisfy the energy conservation in the δ function, $S(\mathbf{q}, \omega)$ must be peaked at $\omega \approx \varepsilon_q$. Actually this leads to a two-peak structure corresponding to $s' = \pm 1$ near $\omega = \varepsilon_q$ for a fixed **q**. But the separation of these two peaks is proportional to $\varepsilon_{(0,0)}$ and is very small at low temperatures. In the numerical calculations performed, the gap between the two peaks is healed by substituting the δ function by a Lorentzian with a small broadening width. As a result of this small broadening, $\mathcal{S}(\mathbf{q},\omega)$ only shows a single-peak structure. Therefore, in this limit the peak positions of $S(\mathbf{q}, \omega)$ follow the boson dispersion.

To better discuss the anisotropic distribution of the spectral weight in momentum space, we plot the constant energy cuts of the calculated $S(\mathbf{q},\omega)$ at a fixed temperature $T < T_{\sigma 0}$ in Fig. 4. At low energies, the peaks of $S(\mathbf{q},\omega)$ form an elliptic ring centered at $(\pi,0)$ [and also its symmetry-related point $(0,\pi)$ after rotation symmetrization], as displayed in Figs. 4(a) and 4(b). The elliptic feature is a consequence of the anisotropic correlation lengths in the Ising-ordered phase, and the ellipticity near $(\pi,0)$ is proportional to $\xi_x/\xi_y = v_{1x}/v_{1y}$, which is not sensitive to temperature dependent for $T < T_{\sigma 0}$ (Fig. 2). The ellipticity also only weakly depends on K: for $J_1/J_2 = 1$, we find $\xi_x/\xi_y \simeq 1.7$ at K = 0 and $\xi_x/\xi_y \simeq 1.4$ at $K/J_2 = 0.8$.

With increasing energy, the ellipse centered around $(\pi, 0)$ expands towards the Brillouin zone boundary, as shown in Figs. 4(a)-4(d). For sufficiently high energy, the spectral weight decreases greatly along the q_x direction, and the $S(\mathbf{q}, \omega)$ is peaked near $q_v = \pm \pi/2$ along the q_v direction [Fig. 4(c)]. The elliptical peak feature appears to have been split into two parts in the direction of its major axis. As the energy gets close to $\varepsilon_{(\pi,\pi)}$, the two peaks move towards $(\pi, \pm \pi)$, forming patterns that are centered around $(\pi, \pm \pi)$ [cf. Figs. 4(c) and 4(d)]. In our theory, there are two factors that contribute to this anisotropic distribution of the spectral weight along the ellipses. First, for $\omega > \varepsilon_{(\pi/2,0)}$, along the q_x axis the energy conservation in the δ function of Eq. (25) can only be satisfied when $\mathbf{k} \neq (0,0)$. A nonzero **k** corresponds to a smaller $n_{\mathbf{k}}$, which greatly reduces $S(q_x, \omega)$. Along the q_y axis, however, $S(q_v,\omega)$ is not reduced because the $\mathbf{k} = (0,0)$ mode can still satisfy the energy conservation. Second, for a given ${\bf k},$ the coherence factor $\cosh(2\theta_{\mathbf{k}} - 2\theta_{\mathbf{k}+\mathbf{q}})$ along the ellipse is also anisotropic. To see this, recall that the largest contribution to



FIG. 4. (Color online) Constant energy cuts of the rotational symmetrized spin dynamical structure factor in the momentum space in the MSW theory for S = 1, $J_1/J_2 = 1.0$, $K/J_2 = 0.8$, and $T/J_2 = 1.0$. The corresponding energies are, respectively, (a) $\omega = 4J_2$, (b) $\omega = 10J_2$, (c) $\omega = 11.5J_2$, and (d) $\omega = 12J_2$. In (a)–(d), a broadening factor $0.5J_2$ has been used for the convenience of calculation.

 $S(\mathbf{q},\omega)$ is from the $\mathbf{k} = (0,0)$ term in Eq. (25). For simplicity, we take a single-mode approximation, namely, $S(\mathbf{q},\omega)$ can be approximated by this $\mathbf{k} = (0,0)$ term. Then the ellipse showing spectral weight peaks is determined by $\varepsilon_q = \omega$, and the coherence factor $\cosh(2\theta_k - 2\theta_{k+q}) \propto (B_q - A_q)/\varepsilon_q$. For $\sqrt{q_x^2\xi_x^2+q_y^2\xi_y^2} \ll T/\Delta_1$, $B_{\mathbf{q}} - A_{\mathbf{q}} \approx \Delta - \varepsilon_{\mathbf{q}}/\Delta - 2\tilde{J}_{1y}f_yq_y^2$, where $\Delta = 8J_2g_{x+y} + 4\tilde{J}_{1x}g_x$. Since $\tilde{J}_{1y} < 0$ for the choice of model parameters, it is easy to see that along the ellipse $\varepsilon_{\mathbf{q}} = \omega$, the maximum of the coherence factor is located along the q_y axis but not the q_x axis. Since within the singlemode approximation, $S(\mathbf{q}, \omega)$ is proportional to the coherence factor, $S(\mathbf{q},\omega)$ is also anisotropic along the ellipse. Note that at low energies $(\sqrt{q_x^2\xi_x^2+q_y^2\xi_y^2} \ll T/\Delta_1), J_{1y}f_yq_y^2 \ll \Delta$, so the anisotropy is very small. This coherence-factor-induced anisotropy becomes sizable when the ellipse is large (for $\sqrt{q_x^2\xi_x^2+q_y^2\xi_y^2}\gtrsim T/\Delta_1$).

V. DISCUSSION

A. Effects of spin size

Besides the S = 1 results shown in Secs. III and IV, we have also studied the J_1 - J_2 -K model with larger spin sizes and found that the mean-field phase diagram is similar to the one in Fig. 1. Approximately, $T_{\sigma 0}$ and T_0 are increased by a factor of S(S + 1)/2. The boson dispersion shown in Fig. 5(a) also exhibits features similar to those in the S = 1 case. In



FIG. 5. (Color online) (a) MSW dispersion along high-symmetry directions in the paramagnetic Brillouin zone for the 2D J_1 - J_2 -K model at S = 2, $J_1/J_2 = 1.0$, $T/J_2 = 1.0$, and various K values. (b) The ratio $\tilde{J}_{1y}/\tilde{J}_{1x}$, showing the anisotropy in effective exchange couplings [see Eq. (9)] as a function of the biquadratic coupling for S = 1 and S = 2.

Fig. 5(b) we compare the ratio of the effective nearest-neighbor couplings $\tilde{J}_{1y}/\tilde{J}_{1x}$, defined in Eq. (9), for S = 1 and S = 2 (at zero temperature). We find that with increasing *S*, the minimal K/J_2 value where \tilde{J}_{1y} becomes ferromagnetic is dropped from $K/J_2 \approx 0.53$ to $K/J_2 \approx 0.13$. Hence, we conclude that the anisotropy of the effective exchange couplings induced by non-Heisenberg coupling *K* is more significant for a larger spin size *S*.

We can further compare our MSW result at T = 0 with the one in a recent MSW study, which used a mean-field treatment that is different from ours.³⁵ The two theories yield exactly the same results when the spin size $S \rightarrow \infty$. For finite spin sizes, by comparing the behavior of the $\tilde{J}_{1y}/\tilde{J}_{1x}$ ratio in Fig. 5(b) and the corresponding results in Ref. 35, we observe that the two theories give qualitatively similar results for the anisotropy in exchange couplings: The biquadratic coupling *K* reduces the effective ratio $\tilde{J}_{1y}/\tilde{J}_{1x}$. Quantitatively, there are some differences between the two approaches. In particular, while for S > 1, the ratio $\tilde{J}_{1y}/\tilde{J}_{1x}$ changes sign at a finite *K* value in both theories, this sign change does not appear for S = 1 in Ref. 35.

B. Generalizations of the J_1 - J_2 -K model

Several remarks on the J_1 - J_2 -K model studied in this paper are in order. From the incipient Mott picture, when the system is in the vicinity of U_c/t , the spin Hamiltonian contains interactions involving more than just two spins. To see this, we start from a multiorbital Hubbard model on a square lattice and assume that Hund's rule coupling locks the spins in different orbitals to a high-spin state. Then we may obtain a spin-only Hamiltonian by integrating out the fermion degrees of freedom based on perturbation in t/U. To the t^2/U order, we obtain the usual J_1 - J_2 Heisenberg interaction between nearest- and next-nearest-neighbor spins. The next-order terms appear in the order t^4/U^3 and include the biquadratic K term as well as the ring exchange interactions. Here, we have focused on the effects of the biquadratic interaction. The influence of the ring exchange interactions in the regime we are considering is briefly discussed in Appendix B.

To fully understand the antiferromagnetic $(\pi, 0, \pi)$ order revealed in the experiments, the 2D J_1 - J_2 -K model needs to be extended to the 3D case by including an interlayer coupling J_z . A nonzero J_z will support the antiferromagnetic order up to the Néel temperature T_N . In the mean-field treatment, the antiferromagnetic order emerges at a mean-field Néel temperature T_{N0} . The details of the effects of the interlayer coupling J_z to the magnetic phase diagram of the J_1 - J_2 -Kmodel and the magnetic excitation spectrum are discussed further in Appendix C.

When fluctuations beyond the mean-field level are taken into account, the actual Néel and Ising transition temperatures, T_N and T_{σ} , can be well below their mean-field values. The mean-field temperatures T_{N0} and $T_{\sigma 0}$ then correspond to some crossover temperature scales, below which the fluctuating order has significant effects. The fluctuating anisotropic effects we have presented will be dominant in the temperature regime $T_N < T < T_{\sigma 0}$.³⁸

C. Effect of itinerant electrons

Within the bad-metal description of the iron pnictides, the quasilocalized moments are coupled to itinerant electrons with a spectral weight that depends on the proximity to the Mott transition. A convenient way to describe the effect of itinerant electrons on the spin dynamics is to reformulate the results of the local-moment-based calculations in terms of a nonlinear σ model and introduce into the latter a damping caused by itinerant electrons; for details, we refer to Ref. 38. Well below $T_{\sigma 0}$ and in the vicinity of (π ,0), the effects of the itinerant electrons are described in terms of the effective action for the staggered magnetization **M**:

$$S[M] = T \int d\mathbf{q} \sum_{l} \left[r + \Delta r + v_{1x}^2 q_x^2 + v_{1y}^2 q_y^2 + \omega_l^2 + \gamma |\omega_l| \right] \mathbf{M}^2 + u \mathbf{M}^4 + \mathcal{O}(\mathbf{M}^6).$$
(26)

Here, $\mathbf{M} = \mathbf{m} + \mathbf{m}'$ is the sum of \mathbf{m} and \mathbf{m}' , the O(3) vectors, respectively, for the magnetizations of the two decoupled sublattices on the square lattice, and ω_l the Matsubara frequency. This action arises in a "*w* expansion," which is based on a proximity to the Mott transition and is described in Refs. 12 and 13; it has the form of the usual σ model.⁵¹ In the first term, $\Delta r > 0$ is a mass shift and γ describes the strength of spin damping from coupling to fermions (see Fig. 6). At relatively low energies, this introduces a procedure that can be used to describe the broadening of the spin spectral peaks in momentum space due to coupling to itinerant electrons.³⁸



FIG. 6. Diagram of the second-order contribution to the effective action in Eq. (26) due to coupling to fermions.

We should emphasize that this procedure is a qualitative treatment of the spin damping. Incorporating the full details of the electronic band structure will introduce momentum dependence of the damping rate, making it possible to generate the type of anisotropic damping that was proposed phenomenologically by Harriger *et al.*³⁹

Comparing our results for the J_1 - J_2 -K model in Figs. 4(a)– 4(d) with those of the J_1 - J_2 model (Fig. 4 in Ref. 38) shows that the biquadratic term itself brings out an anisotropy in the spectral weight of the elliptic peaks. The spectral weight is larger along the major axis of the ellipse than along its minor axis. This anisotropy goes in the same direction as that of the experimental data on BaFe₂As₂, illustrated in Fig. 7(e) and 7(f). We therefore conclude that both the ellipticity and the intensity anisotropy of the spectral peaks in momentum space are controlled by the exchange interactions.

We note that the Ising order parameter is also coupled to the itinerant electrons. Since the Ising order parameter breaks the C_{4v} symmetry, it couples to those spin singlet fermion bilinears that correspond to the B_{1g} representation. Consequently a nonzero Ising order parameter will induce a nonzero $d_{x^2-y^2}$ nematic charge density $(\cos k_x - \cos k_y)c^{\dagger}_{\mathbf{k},\alpha}c_{\mathbf{k},\alpha}$ for all 3*d*orbital electrons, where α is the orbital index. In addition, the Ising order parameter will induce a nonzero charge density imbalance $c_{\mathbf{k}xz}^{\dagger}c_{\mathbf{k}xz} - c_{\mathbf{k}yz}^{\dagger}c_{\mathbf{k}yz}$ between the d_{xz} and the d_{yz} orbitals, which is also referred to as the ferro-orbital order. As a result, the spin fluctuations from the incoherent degrees of freedom can give rise to an orbitally ordered, charge nematic metal, with anisotropic transport properties. In a model with 3D coupling (see Appendix C), the coupling to the itinerant electrons will reduce the Néel transition temperature from its mean-field value T_{N0} to $T_N < T_{N0}$ through the positive Δr noted above. It will likewise decrease the Ising transition temperature from its mean-field value $T_{\sigma 0}$ to $T_{\sigma} < T_{\sigma 0}$. However, the correlation lengths are still sizable and should be anisotropic up to $T_{\sigma 0}$.³⁸ This implies that, in the 3D model with 3D coupling, we expect anisotropic magnetic excitations to exist from T_N all the way up to the crossover temperature scale $T_{\sigma 0}$, in the absence of a static Ising order.

D. Comparison with other approaches

Our studies in the J_1 - J_2 -K model, with or without the coupling to the itinerant electrons, are very different from purely itinerant studies with U/t much smaller than U_c/t . Because the Fermi surface comprises small electron and hole pockets, such calculations are expected to yield very small spin spectral weights. Experimentally, the total spectral weight is known to be large, with an effective moment that is larger than 1 μ_B/Fe in CaFe₂As₂.²¹ Such a large spectral weight arises naturally in our approach using as the starting point the J_1 - J_2 model (with or without the K term).

Our approach can be compared more closely with that of the DMFT studies in Ref. 41, in which the ratio of the effective interaction (combined Coulomb and Hund's interactions) to the characteristic bandwidth is close to the Mott-transition value, $U/t \lesssim U_c/t$. The proximity to the Mott transition ensures that a large part of the electronic spectral weight lies in the incoherent regime, which will naturally give rise to a large spin spectral weight. The consistency of the momentum dependence determined by the DMFT calculations and that of our J_1 - J_2 -K calculations further suggests the compatibility of the two approaches. There are some important differences, however. In the DMFT calculation, the anisotropy of the structure factor has been attributed to the geometry of the Fermi surface(s). The J_1 - J_2 -K results, however, the anisotropy of the spin spectral weight in momentum space with the Ising correlations.

Experimentally, the Ising correlations can be very naturally connected with the *x*-*y* anisotropy observed in ARPES⁵² and transport⁵³ measurements in detwinned 122 iron pnictides at temperatures above T_N . A recent theoretical calculation⁵⁴ shows how resistivity anisotropy in the tetragonal phase above T_N follows from the existence of the Ising correlations discussed here.

VI. COMPARISON WITH EXPERIMENTS ON THE PARAMAGNETIC PHASES OF PARENT 122 IRON PNICTIDES

Spin dynamics in the paramagnetic phase of the parent 122 iron pnictides has recently been studied via INS measurements.^{37,39,40} For CaFe₂As₂, spin dynamics at low energies (below 70 meV) has been studied by Diallo *et al.*³⁷ It was found that the peaks of the dynamical structure factor form anisotropic elliptic features at low energies, similar to the results in the antiferromagnetic phase.²¹ More recently, Harriger *et al.*³⁹ measured the spin dynamics of BaFe₂As₂ up to 200 meV. At low energies, they found that the distribution of spectral weights in the momentum space forms a similar elliptic feature as in the CaFe₂As₂ case. With increasing energy, the elliptic feature expands towards the Brillouin zone boundary. Moreover, they determined the magnetic dispersion to be peaked (or flat-topped) near (π,π). Similar results have also been reported for SrFe₂As₂.⁴⁰

Our study of the J_1 - J_2 -K model has already provided valuable information for understanding these experimental observations. In real materials, the various fluctuation mechanisms and the coupling to fermions/phonons will reduce the Néel and Ising transition temperatures. However, below the mean-field Ising temperature $T_{\sigma 0}$, the effective couplings between the nearest neighbors are always anisotropic. Hence we expect the magnetic fluctuations to be anisotropic for $T_N(\leq T_{\sigma}) < T < T_{\sigma 0}$, which corresponds to the upper portion of region II in Fig. 1. This anisotropy is reflected in the spin dynamics in the paramagnetic phase.

To be specific, the anisotropic elliptic feature at low energies observed in CaFe₂As₂ and other parent 122 compounds can already been understood within the J_1 - J_2 model.³⁸ We show in Fig. 4 that the J_1 - J_2 -K model gives a similar low-energy elliptic feature. It will be important to measure the spin dynamics at high energies in this material.

Our calculated evolution of this elliptic feature as the energy is raised in the J_1 - J_2 -K model can be systematically compared with the experimental observations in $BaFe_2As_2$ and SrFe₂As₂. To see this, we fit the peak positions of the calculated $\mathcal{S}(\mathbf{q},\omega)$ to the experimental magnetic excitation dispersion data in BaFe₂As₂, from which we can extract the best-fitted values of the exchange couplings. Assuming S = 1, we find that the fitted exchange couplings are $J_2 = 17 \pm 4$ meV, $J_1/J_2 = 1.0 \pm 0.5$, and $K/J_2 = 0.8 \pm 0.3$. We find that a very broad range of J_1/J_2 ratio values can all fit the experimental data quite well. As illustrated in Fig. 3(b), any dispersion curve within the shaded region fits the experimental data within error bars. But to fit the dispersion data near the local maximum at (π,π) , a moderate K/J_2 ratio is necessary. For S = 1, we find that $K/J_2 \approx 0.8$ fits the data the best. On the other hand, for S = 2, the best-fitted K/J_2 ratio is substantially reduced, to about 0.2.

For BaFe₂As₂, detailed measurements in momentum space have been reported by Harriger *et al.*³⁹ This allows us to see that the agreement between our theory and the experiment is not only for the dispersion, but also for the anisotropic distribution of the spectral weight of $S(\mathbf{q}, \omega)$ in momentum space.

In order to make a comparison with experimental data, we use Eq. (25) in the calculation of $S(\mathbf{q},\omega)$ and approximate the

 δ function by the following Lorentzian broadening:

$$\delta(\omega - \Delta\varepsilon) \longrightarrow \frac{1}{\pi} \frac{\gamma}{(\omega - \Delta\varepsilon)^2 + \gamma^2}.$$
 (27)

Here we have assumed that the broadening mainly comes from the damping effect due to coupling to itinerant electrons. It is then reasonable to take the phenomenological broadening factor to be the damping γ introduced in Eq. (26) since in either the MSW or the Schwinger boson theory, the damping is still due to the same bubble in Fig. 6. Calculating the magnitude of γ requires a detailed microscopic theory and is beyond the scope of this article; however, we can use Ref. 38 for reference, where it has been determined that $\gamma/J_2 \approx 3$ for CaFe₂As₂. Here we assume that this ratio still holds for BaFe₂As₂ and the damping is isotropic. In Figs. 7(a)-7(d)we replot the theoretical dynamical spin structure factor in Fig. 4 with this damping factor and compare them with the experimental data in Ref. 39. At low energies, our theory correctly captures the elliptical feature centered at $(\pi, 0)$ as displayed in Figs. 7(a) and 7(b). Experimentally, this is seen as a filled elliptical spot due to the damping effect, which is also shown in our theoretical plot in Fig. 7(a). The evolution of the elliptical feature with increasing energy is also consistent with the experimental observation: as the ellipse expands towards



FIG. 7. (Color online) Evolution of $S(\mathbf{q},\omega)$ in the paramagnetic phase of the J_1 - J_2 -K model, showing that the elliptical features near $(\pi, 0)$ at low energies (a),(b) are split into features that are centered around (π, π) , as the energy is increased towards the zone-boundary spin-excitation energy (c),(d). This trend is consistent with the inelastic neutron scattering experiments, shown in the box at the right for two energies measured in the paramagnetic phase of BaFe₂As₂ (data taken from Ref. 39). (a)–(d) Same as Figs. 4(a)–4(d), but with damping $\gamma = 3J_2$. (e),(f) INS data at T = 150 K in BaFe₂As₂, taken from Ref. 39. The energy transfer is (e) $\omega = 50 \pm 10$ meV and (f) $\omega = 150 \pm 10$ meV. Here we find that the best agreement between theory and experimental data is achieved when taking $J_2 \approx 13$ meV in the model.

the zone boundary, it gradually splits into two parts and forms a pattern around (π,π) [see Figs. 7(c), 7(d), and 7(f)]. We reiterate that such anisotropic features are the properties of our J_1 - J_2 -K model either with an isotropic or without additional damping due to itinerant electrons. While the anisotropic damping proposed in Ref. 39 could reinforce the effect, it is not necessary to understand the INS experiments. In CaFe₂As₂ the elliptical feature around $(\pi, 0)$ persists up to high energies, while in BaFe₂As₂ this elliptical feature splits into two parts at intermediate energy.³⁹ These two different behaviors can both be understood within our J_1 - J_2 -K model with similar, nearly isotropic damping, but different K, values.

VII. CONCLUSIONS

In this paper we have investigated the finite-temperature spin dynamics of a J_1 - J_2 -K antiferromagnetic Heisenberg model using both MSW and SBMF theories. The spin dynamics obtained from these two methods are similar to each other.

We have found that by including a moderate biquadratic coupling K, the magnetic excitation spectrum of the J_1 - J_2 -K model is anisotropic below a mean-field Ising transition temperature $T_{\sigma 0}$. As in the case of the J_1 - J_2 model,³⁸ the peak of the low-temperature dynamical structure factor $S(\mathbf{q}, \omega)$ contains elliptical features near $(\pi, 0)$ in the paramagnetic Brillouin zone at low excitation energies. However, unlike the pure J_1 - J_2 model, the spectral intensity also displays anisotropy along the ellipse, with the intensity being higher along the major axis than along the minor axis. This spectral anisotropy accounts for the observed particular way in which the low-energy elliptical features, centered around $(\pi, 0)$, expand towards the zone boundary as the energy is increased towards the zone-boundary spin-excitation energy. It also gives rise to a particular form of high-energy spectral features that are centered around (π,π) .

We have also compared our calculated dynamical spin structure factor of the J_1 - J_2 -K model with the recent inelastic neutron scattering measurements in the paramagnetic phases of 122 iron pnictides.^{37,39,40} The theoretical results provide a very natural understanding of the salient features of the experiments.

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APPENDIX A: ISING TRANSITION AT LOW J_1/J_2 RATIOS

We find that the nature of the mean-field Ising transition at $T_{\sigma 0}$ depends on both the J_1/J_2 and the K/J_2 ratios. At K = 0 and $J_1/J_2 \leq 0.9$, we find $T_{\sigma 0} < T_0$, and the Ising transition at $T_{\sigma 0}$ is always second-order (Fig. 8). When $J_1/J_2 \gtrsim 0.9$, $T_{\sigma 0}$



FIG. 8. (Color online) Mean-field magnetic phase diagram in the MSW theory for S = 1, $J_1/J_2 = 0.6$. The dashed (blue) and dasheddotted (brown) curves refer to the mean-field temperature scales T_{N0} and T_0 , respectively. The thicker solid (red) curve refers to a secondorder Ising transition at $T_{\sigma 0}$, while the thinner solid (red) curve refers to a first-order transition. In the shaded region, the effective exchange coupling $\tilde{J}_{1y} < 0$.

meets T_0 and the Ising transition becomes first-order. This is an artifact of the mean-field approximation since the transition at T_0 is always first-order.^{42,43} Still for $J_1/J_2 \leq 0.9$, increasing *K* from 0, the transition at $T_{\sigma 0}$ changes from second-order to first-order when *K* is bigger than a bicritical point K_c . As shown in Fig. 8 for $J_1/J_2 = 0.6$, $K_c/J_2 \approx 0.04$.

APPENDIX B: EFFECTS OF RING EXCHANGE COUPLINGS

Besides the quadratic and biquadratic interactions, other interactions involving more than two spins can also appear in the spin Hamiltonian in the vicinity of the Mott transition. For instance, a four-spin ring exchange interaction can appear as a consequence of the fourth-order perturbation associated with the electron hopping process. We can consider the effects of a four-spin ring exchange process on the spin dynamics by adding a term $K_{\Box} \sum_{i,j,k,l} [(\mathbf{S}_i \cdot \mathbf{S}_j)(\mathbf{S}_k \cdot \mathbf{S}_l) - (\mathbf{S}_i \cdot \mathbf{S}_k)(\mathbf{S}_j \cdot \mathbf{S}_l) + (\mathbf{S}_i \cdot \mathbf{S}_l)(\mathbf{S}_j \cdot \mathbf{S}_k)]$ to the Hamiltonian, where $K_{\Box} > 0$, and sites (i, j, k, l) are the vertices of a square plaquette, labeled clockwise. The four-spin ring exchange competes against J_1 and J_2 and tends to weaken the antiferromagnetic order coming from J_1 or J_2 . In the linear spin-wave description of the $(\pi,0)$ ordered state, we obtain the effective exchange constants $\tilde{J}_{1x} = J_1 + 2(K - K_{\Box})S^2$, $\tilde{J}_{1y} = J_1 - 2(K - K_{\Box})S^2$, and $\tilde{J}_2 = J_2 + K_{\Box}S^2$ and the reduced spin gap at (π, π) . This trend also persists in the paramagnetic state and reduces the size of the Ising order parameter. For consistency with the experimental results we require $K > K_{\Box}$.

APPENDIX C: EFFECTS OF INTERLAYER EXCHANGE COUPLING

The real materials have a 3D tetragonal structure. In the J_1 - J_2 -K model, the 3D effects can be studied by extending the model to include a finite interlayer exchange interaction



FIG. 9. (Color online) Mean-field magnetic phase diagram in the MSW theory for S = 1, $J_1/J_2 = 1$, and an interlayer exchange coupling $J_z/J_2 = 0.1$. The dashed (blue) and solid (red) curves refer to the mean-field temperature scales T_{N0} and $T_{\sigma 0}$, respectively. In the shaded region, the effective exchange coupling $\tilde{J}_{1y} < 0$.

 $J_z \sum_i \mathbf{S}_i \cdot \mathbf{S}_{i+\hat{z}}$. In three dimensions the long-range antiferromagnetic phase survives at finite temperatures up to the Néel temperature T_N . In the MSW and SBMF theories, the mean-field Néel temperature T_{N0} is determined by the onset of spontaneous sublattice magnetization m_0 . In general, $T_{N0} \leq T_{\sigma 0} \leq T_0$. The modification to our discussion in Sec. II comes through an additional interlayer antiferromanetic bond correlation parameter g_z . In the presence of J_z , the selfconsistent equations (4)–(6) and (18)–(20) are unchanged, but the expressions for A_k and B_k are modified according to

$$A_{\mathbf{k}}^{\mathrm{3D}} = A_{\mathbf{k}} + 2J_z g_z \cos k_z, \qquad (C1)$$

$$B_{\mathbf{k}}^{\mathrm{3D}} = B_{\mathbf{k}} + 2J_z g_z, \tag{C2}$$

in the MSW mean-field theory and

$$A_{\mathbf{k}}^{3\mathrm{D}} = A_{\mathbf{k}} + 2J_z g_z \sin k_z, \qquad (C3)$$

$$B_{\mathbf{k}}^{\mathrm{3D}} = B_{\mathbf{k}}.$$
 (C4)

in the SBMF theory.

In Fig. 9 we show the phase diagram at the experimentally suggested ratio $J_z/J_2 = 0.1$. Similarly to the 2D case, the mean-field phase diagram consists of an Ising- and Néel-ordered antiferromagnetic phase (I), an Ising-ordered but Néel-disordered paramagnetic phase (II), and an Ising- and Néel-disordered paramagnetic phase (III), separated by mean-field temperatures T_{N0} and $T_{\sigma0}$ (see also Fig. 10). For the parameters in Fig. 9, the transitions are both first-order, and both T_{N0} and $T_{\sigma0}$ increase with *K*. For $K/J_2 \gtrsim 0.2$, T_{N0} meets $T_{\sigma0}$, and there is only a single transition between phase I and phase III. The absence of phase II in this regime is an artifact of the mean-field theory, since $T_{\sigma0}$ is always bounded above by the mean-field scale T_0 .

In connection to the real materials, we note that the structural and magnetic transitions in the 1111 pnictides are well separated. But in 122 compounds, either they are very close to each other or they become a single first-order transition. This can be understood in terms of the present theory, provided J_z is stronger in the 122 materials.



FIG. 10. (Color online) Temperature evolution of the mean-field parameters in the MSW theory for S = 1, $J_1/J_2 = 1$, and $K/J_2 = 0.8$, with an interlayer exchange coupling $J_z/J_2 = 0.1$.

By comparing Figs. 9 and 2 we see that the magnetic transition is closer to the Ising transition for a larger J_z . Recent experiments also show that electron doping may cause separation of the structural and magnetic transition temperatures in the Ba(Fe,Co)₂As₂ system.⁵⁵ The similarity between this behavior and the *K* dependence of $T_{\sigma 0}$ and T_{N0} in the phase diagram in Fig. 9 suggests the possibility that electron doping is positively correlated with a reduction of the biquadratic interaction. It would then be interesting to reveal the link between them in future experimental and theoretical studies.

In Fig. 11 we show the low-temperature boson dispersions of the 3D model for various K values along two high-symmetry directions in the $k_z = \pi$ plane. Aside from a larger gap at



FIG. 11. (Color online) Dispersion of the J_1 - J_2 -K model in the MSW theory for various K values at S = 1, $J_1/J_2 = 1.0$, and $T/J_2 = 0.1$, with an interlayer exchange coupling $J_z/J_2 = 0.1$.

 $(0,0,\pi)$, the dispersion is very similar to the one in 2D: the dispersion is highly anisotropic, and with increasing *K*, the local minimum at (π,π,π) turns to a maximum. This is not too

surprising because the in-plane anisotropy is a consequence of the 2D Ising-type fluctuations and is not sensitive to the interlayer exchange coupling.

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