

Electronic orders in multiorbital Hubbard models with lifted orbital degeneracyShintaro Hoshino¹ and Philipp Werner²¹*Department of Basic Science, The University of Tokyo, Meguro, Tokyo 153-8902, Japan*²*Department of Physics, University of Fribourg, 1700 Fribourg, Switzerland*

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We study the symmetry-broken phases in two- and three-orbital Hubbard models with lifted orbital degeneracy using dynamical mean-field theory. On the technical level, we explain how symmetry relations can be exploited to measure the four-point correlation functions needed for the calculation of the lattice susceptibilities. In the half-filled two-orbital model with crystal-field splitting, we find an instability of the metallic phase to spin-orbital order with neither spin nor orbital moment. This ordered phase is shown to be related to the recently discovered fluctuating-moment induced spin-triplet superconducting state in the orbitally degenerate model with shifted chemical potential. In the three-orbital case, we consider the effect of a crystal-field splitting on the spin-triplet superconducting state in the model with positive Hund coupling, and the spin-singlet superconducting state in the case of negative Hund coupling. It is demonstrated that for certain crystal-field splittings the higher energy orbitals instead of the lower ones are relevant for superconductivity, and that T_c can be slightly enhanced by the crystal-field effect. We comment on the implications of our results for the superconductivity in strontium ruthenates, and for the recently reported light-enhanced superconducting state in alkali-metal-doped fullerenes.

DOI: [10.1103/PhysRevB.93.155161](https://doi.org/10.1103/PhysRevB.93.155161)**I. INTRODUCTION**

The local Slater-Kanamori interaction [1,2] originating from Coulomb repulsion leads to highly nontrivial phase diagrams and crossover phenomena in multiorbital Hubbard models. Depending on the filling and the energy splittings between the orbitals one finds antiferromagnetic or ferromagnetic order [3,4], orbital order [3], high-spin/low-spin transitions [5,6], staggered high-spin/low-spin order [7], excitonic insulating phases [8,9], or intraorbital spin-triplet (equal-spin) pairing [4]. Some of these instabilities are linked to the spin-freezing crossover [10], which occurs in models with nonzero (positive) Hund coupling, and underlies the unusual finite-temperature properties of Hund metals [11]. In models with negative Hund coupling, an intraorbital spin-singlet superconducting state appears [12,13], but alternative ordered states have not yet been systematically explored.

An unbiased way of mapping out the electronic instabilities to long-range orders is to compute the corresponding susceptibilities and to look for divergences as a function of the model parameters. While such calculations cannot be easily performed with numerically exact lattice methods, they become computationally tractable within the dynamical mean field theory (DMFT) approximation [14]. This theory assumes a spatially local self-energy and vertex, and produces qualitatively correct solutions for high-dimensional lattice models. The use of a featureless density of states in DMFT ensures that the results do not depend on subtle band-structure effects. In this work, we employ the DMFT formalism and a semicircular density of states to map out the instabilities to uniform and staggered long-range-ordered phases in two and three orbital systems, focusing on the physically most interesting intermediate coupling regime.

One main purpose of this study is to show how symmetry relations can be exploited in the numerical simulations based on continuous-time Monte Carlo impurity solvers [15], and in the analysis of the ordering instabilities. In particular, we will demonstrate that the spin-orbital ordered phase (also

called excitonic insulator phase [8,9]) appearing in half-filled two-orbital models with crystal-field (CF) splitting is related by symmetry operations to the orbital-singlet spin-triplet superconducting state found in the orbitally degenerate model away from half filling [16–21]. We also use symmetry relations to analyze and explain the Cooper pair formation in three orbital systems with lifted orbital degeneracy.

For positive (ferromagnetic) Hund coupling, the Slater-Kanamori interaction describes t_{2g} -based three orbital systems, as realized for example in Sr_2RuO_4 and SrRuO_3 [11]. These materials indeed exhibit spin-triplet superconductivity or ferromagnetism, as well as the non-Fermi-liquid properties associated with spin freezing. Recently the authors have demonstrated that a spin-triplet pairing is induced by the fluctuating local moments which appear at the border of the spin-frozen regime and eventually order ferromagnetically or antiferromagnetically [4]. Similar multiorbital physics is also expected to be relevant for other types of spin-triplet pairing, such as realized in U-based ferromagnetic superconductors [22]. In this paper we will examine the effect of CF splittings on the spin-triplet pairing. We will show that the Cooper pairs can be formed in the higher-energy orbitals instead of the lower one. This somewhat counterintuitive result can be explained by exploiting a particle-hole transformation.

The half-filled three-orbital model with negative Hund coupling exhibits an intraorbital pairing state which is relevant for fulleride superconductors [12,23,24]. The effect of CF splittings on this superconducting state is of interest in connection with the recently reported light-enhanced superconductivity in K_3C_{60} [25]. We show that in equilibrium, the lifting of the orbital degeneracy can slightly stabilize the pairing, but the effect on T_c is small. Hence, it is unlikely that the experimentally observed dramatic enhancement of T_c can be explained by quasistatic distortions of the C_{60} molecules within an equilibrium picture.

The outline of the paper is as follows. In Sec. II, we present the model and explain its basic properties, in particular the symmetries. We discuss in Sec. III some technical details

concerning the calculation of lattice susceptibilities within DMFT. Section IV shows the results for the two-orbital model, and explains the connection between spin-orbital order and spin-triplet superconductivity, while Sec. V presents results for the three-orbital model with lifted orbital degeneracy. Section VI is a brief summary and conclusion.

II. MODEL AND METHOD

We consider the Hubbard model with M degenerate orbitals given by the general Hamiltonian

$$\begin{aligned} \mathcal{H} = & \sum_{k\gamma\sigma} (\varepsilon_k - \mu) c_{k\gamma\sigma}^\dagger c_{k\gamma\sigma} + U \sum_{i\gamma} n_{i\gamma\uparrow} n_{i\gamma\downarrow} \\ & + U' \sum_{i\sigma, \gamma < \gamma'} n_{i\gamma\sigma} n_{i\gamma'\sigma} + (U' - J) \sum_{i\sigma, \gamma < \gamma'} n_{i\gamma\sigma} n_{i\gamma'\sigma} \\ & - \alpha J \sum_{i, \gamma < \gamma'} (c_{i\gamma\uparrow}^\dagger c_{i\gamma\downarrow} c_{i\gamma'\downarrow}^\dagger c_{i\gamma'\uparrow} \\ & + c_{i\gamma\uparrow}^\dagger c_{i\gamma\downarrow}^\dagger c_{i\gamma'\uparrow} c_{i\gamma'\downarrow} + \text{H.c.}). \end{aligned} \quad (1)$$

The operator $c_{i\gamma\sigma}$ annihilates the electron with orbital γ and spin σ at site i . The Fourier transformation is defined by $c_{k\gamma\sigma} = N^{-1/2} \sum_i c_{i\gamma\sigma} e^{ik \cdot R_i}$, where N is the number of sites and R_i is the spatial coordinate at site i . The number operator is defined by $n_{i\gamma\sigma} = c_{i\gamma\sigma}^\dagger c_{i\gamma\sigma}$. In Eq. (1), the parameter α is introduced to describe the effect of spin anisotropy and orbital anisotropy. Physically this anisotropy originates from the spin-orbit coupling. We note that for repulsively interacting systems, the pair hopping term is irrelevant. This is because the strongest intra-orbital repulsion U disfavors the doubly occupied orbital state and as a result the pair hopping process between orbitals rarely occurs. If the Hund coupling J is negative, as in models discussed in connection with fulleride superconductors [12,24], the pair hopping becomes relevant, while spin-flip processes are effectively suppressed. With an anisotropy parameter $\alpha < 1$, the effect of these spin-flip and pair hopping processes becomes weaker.

Let us first consider the isotropic case with $\alpha = 1$. In rotationally invariant systems, where the relation $U = U' + 2J$ holds, the interaction part of the Hamiltonian can be rewritten in the form

$$\begin{aligned} \mathcal{H}_{\text{int}} = & \sum_{i\gamma\gamma'\sigma\sigma'} \left[\frac{U'}{2} c_{i\gamma\sigma}^\dagger c_{i\gamma\sigma} c_{i\gamma'\sigma'}^\dagger c_{i\gamma'\sigma'} + \frac{J}{2} c_{i\gamma\sigma}^\dagger c_{i\gamma'\sigma} c_{i\gamma'\sigma'}^\dagger c_{i\gamma\sigma'} \right. \\ & \left. + \frac{J}{2} c_{i\gamma\sigma}^\dagger c_{i\gamma'\sigma} c_{i\gamma\sigma'}^\dagger c_{i\gamma'\sigma'} \right], \end{aligned} \quad (2)$$

which is suitable for discussing the symmetries of this model. Using this expression, we can easily verify that the transformation

$$\mathcal{S} c_{i\gamma\sigma} \mathcal{S}^{-1} = e^{i\theta} \sum_{\sigma'} \mathcal{U}_{\sigma\sigma'} \sum_{\gamma'} \mathcal{V}_{\gamma\gamma'} c_{i\gamma'\sigma'} \quad (3)$$

does not change \mathcal{H}_{int} , if \mathcal{U} and \mathcal{V} are 2×2 unitary and $M \times M$ orthogonal matrices, respectively. This invariance implies a $U(1) \times SU(2) \times SO(M)$ symmetry.

On the other hand, in the anisotropic case with $\alpha = 0$, there are only density-density type interactions. Then the symmetry is described by $[U(1)]^{2M}$ where $2M$ is the number of

spin/orbital indices. The relevant symmetry operation is given by $\mathcal{S} c_{i\gamma\sigma} \mathcal{S}^{-1} = e^{i\theta_{\gamma\sigma}} c_{i\gamma\sigma}$ for any (γ, σ) .

When we consider the degenerate-orbital model including kinetic terms, the total Hamiltonian is also unchanged by this local transformation. In this case we have the relation

$$\langle \mathcal{O} \rangle = \langle \mathcal{S} \mathcal{O} \mathcal{S}^{-1} \rangle \quad (4)$$

for arbitrary operators \mathcal{O} . This relation will be used in Sec. III C to derive measurement formulas for the two-particle Green functions.

We analyze the multiorbital Hubbard model within the framework of DMFT [14]. This theory becomes exact in the limit of infinite dimensions, where only local correlations are relevant. Because of the local self-energy, the lattice problem can be mapped onto a multiorbital impurity problem with local interaction terms identical to those of Eq. (1) and noninteracting baths whose properties can be encoded by the hybridization functions $\Delta_{\gamma\sigma}$. We consider the infinite-dimensional Bethe lattice, whose noninteracting density of states has a semicircular shape: $\rho(\varepsilon) = (1/2\pi t^2) \sqrt{4t^2 - \varepsilon^2}$ with $t = 1$ the (rescaled) hopping integral. For this lattice, the DMFT self-consistency condition simplifies to $\Delta_{\gamma\sigma} = t^2 G_{\gamma\sigma}$, with $G_{\gamma\sigma}$ the impurity Green function. Since the wave vector in the Bethe lattice is ill defined, it is natural to work in a real-space representation. On the other hand we can also consider an alternative description in terms of “pseudo-wave-vectors” (see Appendix A), which enables a discussion analogous to that for ordinary periodic lattices.

III. EVALUATION OF SUSCEPTIBILITIES

A. Definition

In this paper we discuss the instabilities toward long-ranged orders. We consider ordered states corresponding to operators

$$\mathcal{O}(\mathbf{q}) = \sum_i \mathcal{O}_i e^{-i\mathbf{q} \cdot \mathbf{R}_i}, \quad (5)$$

which can be detected by the divergence of the susceptibilities defined by

$$\chi_{\mathcal{O}}(\mathbf{q}) = \frac{1}{N} \int_0^\beta \langle \mathcal{O}(\mathbf{q}, \tau) \mathcal{O}(-\mathbf{q}) \rangle d\tau. \quad (6)$$

Here $\mathcal{O}(\tau) = e^{\tau \mathcal{H}} \mathcal{O} e^{-\tau \mathcal{H}}$ is the Heisenberg picture with imaginary time τ . In DMFT, these susceptibilities can be calculated from the local vertex parts extracted from the effective impurity problem [14].

Let us consider the specific forms of \mathcal{O} by taking the two-orbital model as an example. For diagonal orders, the operators \mathcal{O}_i are given by

$$n_i = \sum_{\gamma\sigma} c_{i\gamma\sigma}^\dagger c_{i\gamma\sigma}, \quad (7)$$

$$s_i^\mu = \sum_{\gamma\sigma\sigma'} c_{i\gamma\sigma}^\dagger \sigma_{\sigma\sigma'}^\mu c_{i\gamma\sigma'}, \quad (8)$$

$$\tau_i^v = \sum_{\gamma\gamma'\sigma} c_{i\gamma\sigma}^\dagger \sigma_{\gamma\gamma'}^v c_{i\gamma'\sigma}, \quad (9)$$

$$o_i^{v\mu} = \sum_{\gamma\gamma'\sigma\sigma'} c_{i\gamma\sigma}^\dagger \sigma_{\gamma\gamma'}^v \sigma_{\sigma\sigma'}^\mu c_{i\gamma'\sigma'}, \quad (10)$$

corresponding to charge, spin, orbital, and spin-orbital moments, respectively. Here $\mu, \nu = x, y, z$. For off-diagonal orders, we have

$$p_i^{s\mu} = \frac{1}{2} \sum_{\gamma\gamma'\sigma\sigma'} c_{i\gamma\sigma}^\dagger \epsilon_{\gamma\gamma'} (\sigma^\mu \epsilon)_{\sigma\sigma'} c_{i\gamma'\sigma'}^\dagger + \text{H.c.}, \quad (11)$$

$$p_i^{vs} = \frac{1}{2} \sum_{\gamma\gamma'\sigma\sigma'} c_{i\gamma\sigma}^\dagger (\sigma^\nu \epsilon)_{\gamma\gamma'} \epsilon_{\sigma\sigma'} c_{i\gamma'\sigma'}^\dagger + \text{H.c.}, \quad (12)$$

which correspond to the orbital-singlet-spin-triplet and orbital-triplet-spin-singlet pairing amplitudes, respectively. The anti-symmetric unit tensor is defined by $\epsilon = i\sigma^y$. All of these operators are Hermitian.

To further classify the diagonal operators given above, we consider the time-reversal operation defined by $\mathcal{T} = \exp(-i\pi \sum_i s_i^y/2) \mathcal{K}$ with complex conjugation operator \mathcal{K} . This transforms the electron operator as

$$\mathcal{T} c_{i\gamma\sigma} \mathcal{T}^{-1} = \sum_{\sigma'} \epsilon_{\sigma\sigma'} c_{i\gamma\sigma'}. \quad (13)$$

Note that \mathcal{T} is an antiunitary operator: $\mathcal{T} z \mathcal{T}^{-1} = z^*$ for a complex number z . This operation changes the momentum ($\mathbf{k} \rightarrow -\mathbf{k}$) and flips the spin state ($\uparrow \rightarrow \downarrow$ or $\downarrow \rightarrow \uparrow$). As expected, the charge is time-reversal even ($\mathcal{T} n_i \mathcal{T}^{-1} = n_i$) and spin is odd ($\mathcal{T} s_i^\mu \mathcal{T}^{-1} = -s_i^\mu$). The orbital moments τ_i^x and τ_i^z are time-reversal even, while τ_i^y is odd. The difference between the orbital moments arises due to the presence of the imaginary unit in the Pauli matrix σ^y , which gives an additional minus sign under the time-reversal operation.

The results are summarized in Table I. While in this paper we focus on a model study, a physical interpretation in terms of the doubly degenerate e_g orbitals is given in Appendix B, where the operator τ^y is shown to be a (generalized) magnetic moment. These arguments can also be applied to the three orbital model, where the Pauli matrix for the orbital is replaced by the Gell-Mann matrix. In this case we have eight kinds of orbital orders, three of which are time-reversal odd operators.

For the pairing state characterized by Eqs. (11) and (12), the time-reversal symmetry is not broken. More specifically, while the time-reversal operation \mathcal{T} can change the sign of these operators, this can be absorbed by simultaneously performing a global U(1) gauge transformation. On the other hand, if two or more Cooper pair amplitudes become finite with different phases, then the resulting state has broken time-reversal

TABLE I. Classification of operators relevant to diagonal orders for the rotationally invariant case ($\alpha = 1$).

Identifier	Operator	Time reversal
(i)	n	even
(ii)	s^x, s^y, s^z	odd
(iii)	τ^x, τ^z	even
(iv)	τ^y	odd
(v)	$o^{xx}, o^{xy}, o^{xz}, o^{zx}, o^{zy}, o^{zz}$	odd
(vi)	o^{yx}, o^{yy}, o^{yz}	even
(vii)	p^{sx}, p^{sy}, p^{sz}	even
(viii)	p^{xs}, p^{zs}	even
(ix)	p^{ys}	even

symmetry [26]. In this case the sign cannot be absorbed by the gauge transformation, and thus the time-reversal symmetry is broken.

B. Two-particle Green function and susceptibilities

Here we discuss the calculation of the susceptibility in the framework of DMFT [14]. For diagonal and off-diagonal orders, the relevant two-particle Green functions are given by

$$\begin{aligned} \chi_{ij,a_1a_2a_3a_4}^{\text{diag}}(\tau_1, \tau_2, \tau_3, \tau_4) \\ = \langle T_\tau c_{ia_1}^\dagger(\tau_1) c_{ia_2}(\tau_2) c_{ja_3}^\dagger(\tau_3) c_{ja_4}(\tau_4) \rangle \\ - \langle T_\tau c_{ia_1}^\dagger(\tau_1) c_{ia_2}(\tau_2) \rangle \langle T_\tau c_{ja_3}^\dagger(\tau_3) c_{ja_4}(\tau_4) \rangle, \end{aligned} \quad (14)$$

$$\begin{aligned} \chi_{ij,a_1a_2a_3a_4}^{\text{off-d}}(\tau_1, \tau_2, \tau_3, \tau_4) \\ = \langle T_\tau c_{ia_1}^\dagger(\tau_1) c_{ia_2}^\dagger(\tau_2) c_{ja_3}(\tau_3) c_{ja_4}(\tau_4) \rangle, \end{aligned} \quad (15)$$

respectively. The index a denotes the spin and orbital indices (γ, σ). Here T_τ represents the imaginary time-ordering operator, and we subtract the disconnected part in Eq. (14). We also define the Fourier transformation with respect to imaginary time by

$$\begin{aligned} \chi_{ij,a_1a_2a_3a_4}^\xi(i\varepsilon_n, i\varepsilon_{n'}) \\ = \frac{1}{\beta^2} \int_0^\beta d\tau_1 d\tau_2 d\tau_3 d\tau_4 \chi_{ij,a_1a_2a_3a_4}^\xi(\tau_1, \tau_2, \tau_3, \tau_4) \\ \times e^{i\varepsilon_n(\tau_2-\tau_1)} e^{i\varepsilon_{n'}(\tau_4-\tau_3)}, \end{aligned} \quad (16)$$

where ξ means “diag” or “off-d.” Since we are interested only in static susceptibilities, we set the bosonic frequency to zero. The susceptibilities defined in Eq. (6) can be calculated by summing up the fermionic Matsubara frequencies in the two-particle Green function, which is explicitly written in the form

$$\begin{aligned} \chi_\eta^\xi(\mathbf{q}) = \frac{T}{N} \sum_{nn'} \sum_{ij} \sum_{a_1a_2a_3a_4} \mathcal{A}_{a_1a_2}^\eta (\mathcal{A}_{a_3a_4}^\eta)^* e^{-i\mathbf{q} \cdot (\mathbf{R}_i - \mathbf{R}_j)} \\ \times \chi_{ij,a_1a_2a_3a_4}^\xi(i\varepsilon_n, i\varepsilon_{n'}), \end{aligned} \quad (17)$$

where the $\mathcal{A}_{aa'}$ are form factors originating from Eqs. (7)–(10) for $\xi = \text{“diag”}$ and from Eqs. (11) and (12) for $\xi = \text{“off-d.”}$

Now we employ the Bethe-Salpeter equation which relates the two-particle Green function and vertex. Since the vertex part Γ is local in DMFT [14], it can be evaluated from the local two-particle Green function as

$$\begin{aligned} \chi_{ii,a_1a_2a_3a_4}^\xi(i\varepsilon_n, i\varepsilon_{n'}) \\ = \chi_{ii,a_1a_2a_3a_4}^{\xi 0}(i\varepsilon_n, i\varepsilon_{n'}) + \sum_{n_1n_2} \sum_{aa'a''} \chi_{ii,a_1a_2aa'}^{\xi 0}(i\varepsilon_n, i\varepsilon_{n_1}) \\ \times \Gamma_{i,a'a''a'''}(i\varepsilon_{n_1}, i\varepsilon_{n_2}) \chi_{ii,a''a''a_3a_4}^\xi(i\varepsilon_{n_2}, i\varepsilon_{n'}). \end{aligned} \quad (18)$$

Here the two-particle Green function without vertex parts is written as $\chi^{\xi 0}$. The site index for the vertex part can be neglected if the system is uniform.

The local vertex extracted above is inserted into the nonlocal Bethe-Salpeter equation

$$\begin{aligned} \chi_{ij,a_1a_2a_3a_4}^{\xi}(i\varepsilon_n, i\varepsilon_{n'}) \\ = \chi_{ij,a_1a_2a_3a_4}^{\xi 0}(i\varepsilon_n, i\varepsilon_{n'}) + \sum_{\ell n_1 n_2} \sum_{aa'a''} \chi_{i\ell,a_1a_2aa'}^{\xi 0}(i\varepsilon_n, i\varepsilon_{n_1}) \\ \times \Gamma_{\ell,a'a''a'''}(i\varepsilon_{n_1}, i\varepsilon_{n_2}) \chi_{\ell j,a''a'''a_3a_4}^{\xi}(i\varepsilon_{n_2}, i\varepsilon_{n'}). \end{aligned} \quad (19)$$

In practice, it is convenient to perform the Fourier transformation with respect to the site index before solving the matrix equation. Thus we obtain the two-particle lattice Green functions, which contain the information of the susceptibilities.

C. Measurement trick for local two-particle Green functions

As explained in the previous section, the local vertices can be extracted from the local two-particle Green functions. In DMFT calculations of the multiorbital Hubbard model (1), we must consider the following two-particle Green functions of the corresponding impurity model:

$$\chi_{\gamma\gamma'\sigma\sigma'}^a = \langle T_{\tau} c_{\gamma\sigma}^{\dagger}(\tau_1) c_{\gamma\sigma}(\tau_2) c_{\gamma'\sigma'}^{\dagger}(\tau_3) c_{\gamma'\sigma'}(\tau_4) \rangle, \quad (20)$$

$$\chi_{\gamma\gamma'\sigma\sigma'}^b = \langle T_{\tau} c_{\gamma\sigma}^{\dagger}(\tau_1) c_{\gamma\sigma}(\tau_2) c_{\gamma'\sigma'}^{\dagger}(\tau_3) c_{\gamma'\sigma'}(\tau_4) \rangle, \quad (21)$$

$$\chi_{\gamma\gamma'\sigma\sigma'}^c = \langle T_{\tau} c_{\gamma\sigma}^{\dagger}(\tau_1) c_{\gamma'\sigma}(\tau_2) c_{\gamma\sigma}^{\dagger}(\tau_3) c_{\gamma'\sigma'}(\tau_4) \rangle. \quad (22)$$

All of the diagonal and off-diagonal two-particle Green functions that appear in the previous subsection can be calculated from $\chi^{a,b,c}$. A powerful method to solve impurity models is the continuous-time Monte Carlo technique [15]. Here, we employ the hybridization expansion method in the matrix formulation [27]. In this approach, the function χ^a can be calculated by the standard procedure which involves the removal of two hybridization functions [15]. However, the other functions $\chi^{b,c}$, which are in general finite, cannot be obtained by the standard technique unless the impurity model has off-diagonal hybridizations.

If we have the continuous $SU(2) \times SO(M)$ symmetry the two-particle Green functions χ^b and χ^c can be expressed in terms of χ^a using Eq. (4). To see this, we consider the function $\chi_{12\uparrow\downarrow}^b$ in the two-orbital model ($M = 2$). The $SU(2)$ and $SO(2)$ matrices in the spin and orbital spaces are given by

$$\mathcal{U} = \begin{pmatrix} z & -w^* \\ w & z^* \end{pmatrix}, \quad \mathcal{V} = \begin{pmatrix} x & -y \\ y & x \end{pmatrix}, \quad (23)$$

where $|z|^2 + |w|^2 = 1$ ($z, w \in \mathbb{C}$) and $x^2 + y^2 = 1$ ($x, y \in \mathbb{R}$). The $SO(2)$ matrix in orbital space represents a rotation around the τ^y axis. The quantity $\chi_{12\uparrow\downarrow}^b$ is transformed by \mathcal{U} as

$$\chi_{12\uparrow\downarrow}^b \longrightarrow (|z|^4 + |w|^4) \chi_{12\uparrow\downarrow}^b + 2|z|^2 |w|^2 (\chi_{12\uparrow\uparrow}^a - \chi_{12\downarrow\downarrow}^a), \quad (24)$$

where we have used the equivalence between $\gamma = 1, 2$ and between $\sigma = \uparrow, \downarrow$. If the transformation does not change the whole Hamiltonian, which implies $\alpha = 1$, the left- and right-hand sides of Eq. (24) can be regarded as identical. Thus we obtain the relation

$$\chi_{12\uparrow\downarrow}^b = \chi_{12\uparrow\uparrow}^a - \chi_{12\downarrow\downarrow}^a. \quad (25)$$

Note that this result is not dependent on the values of z and w . This relation has been previously used in the study of a two-channel Kondo lattice [28]. In a similar manner, if the system has orbital $SO(2)$ symmetry, it satisfies the additional relation

$$\chi_{12\uparrow\downarrow}^c = \chi_{11\uparrow\downarrow}^a - \chi_{12\uparrow\downarrow}^a + \tilde{\chi}_{12\uparrow\downarrow}^b, \quad (26)$$

where we have defined the time-sorted quantity

$$\tilde{\chi}(\tau_1, \tau_2, \tau_3, \tau_4) = -\chi(\tau_1, \tau_4, \tau_3, \tau_2). \quad (27)$$

By an analogous trick we can also obtain the relation

$$\chi_{12\uparrow\uparrow}^c = \chi_{11\uparrow\uparrow}^a - \chi_{12\uparrow\uparrow}^a + \tilde{\chi}_{12\uparrow\uparrow}^b. \quad (28)$$

The other types of χ^b and χ^c can be represented by the quantities obtained above. Thus, for the calculation of the two-particle Green functions $\chi^{a,b,c}$, it is sufficient to measure χ^a .

A key prerequisite for this technique is the $SU(2)$ or $SO(2)$ symmetry in the Hamiltonian. Hence, these formulas do not hold when the symmetry is lowered by, e.g., a CF splitting. The generation of new terms on the right-hand side of Eq. (24) is also an important ingredient for the present trick. For example the $U(1)$ transformation only changes the phase factor, which is not enough to derive expressions for χ^b and χ^c . Thus, at least a two-dimensional representation of the symmetry operation is necessary.

Let us also add a comment on the phase transformation. While usually it changes the phase of the fermion operators uniformly ($c_{i\gamma\sigma} \rightarrow c_{i\gamma\sigma} e^{i\theta}$), here we can slightly generalize it as $c_{i\gamma\sigma} \rightarrow c_{i\gamma\sigma} e^{i(\theta_{\sigma} + \theta_{\gamma})}$ because of the $SU(2) \times SO(2)$ symmetry. For the invariance of the Hamiltonian, the phases θ_{\uparrow} and θ_{\downarrow} are arbitrary, but $\theta_1 - \theta_2 = \pi m$ ($m \in \mathbb{Z}$) must be satisfied due to the pair hopping term. These are related to conservation laws of spin/orbital indices. Combining this transformation with Eq. (4), it can be explicitly shown that expectation values such as $\langle c_{1\uparrow}^{\dagger} c_{1\downarrow} c_{1\uparrow}^{\dagger} c_{1\downarrow} \rangle$ and $\langle c_{1\uparrow}^{\dagger} c_{1\uparrow} c_{1\uparrow}^{\dagger} c_{2\uparrow} \rangle$ are identically zero.

The above techniques can also be used in models with a general number M of orbitals. Namely, if we choose two of the M orbitals, we can utilize the partial $SO(2)$ symmetry within this subspace, so that the procedure remains unchanged.

IV. TWO-ORBITAL MODEL

A. Degenerate orbitals

In this section, we study the ordering instabilities in the two-orbital model with $J > 0$ and first consider the orbitally degenerate case [Eq. (1)]. Figure 1 shows the staggered susceptibilities characterized by the “ordering vector \mathbf{Q} ” (see Appendix A) in the spin-isotropic system with $\alpha = 1$. Here the symmetry tricks explained in the previous section are used for the evaluation of the two-particle Green functions. The uniform susceptibilities for diagonal and off-diagonal orders do not diverge for the chosen parameters, and therefore are not shown.

Near half filling ($n \lesssim 2$), the antiferromagnetic order with $s(\mathbf{Q})$ appears as demonstrated by the sign change of the corresponding inverse susceptibility in Fig. 1. The other susceptibilities remain positive and hence no further instabilities are present. For the low-filling case with $n \lesssim 1$,

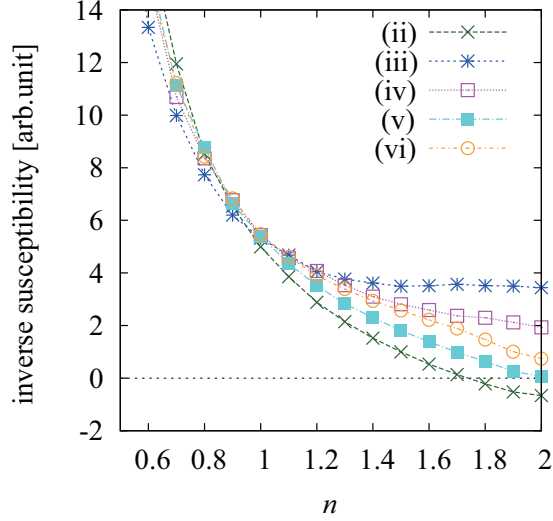


FIG. 1. Filling dependence of inverse staggered susceptibilities for diagonal orderings in the isotropic case ($\alpha = 1$). The orders (ii)–(vi) are defined in Table I. We choose the parameters $U = 2$, $J/U = 1/4$, and $T = 0.02$. The staggered charge susceptibility [(i) in Table I] is not shown here because of low numerical accuracy.

all of the susceptibilities behave in a similar manner. This is because the interaction effect is weaker and an approximate SU(4) symmetry appears. For the present choice of parameters no other symmetry breaking, including superconductivity, occurs.

In our study, although the survey is limited due to the high numerical cost for $\alpha \neq 0$, we did not find any interesting ordered states for the spin isotropic case ($\alpha = 1$) with $J > 0$. In the following, we therefore focus on the model with anisotropic interaction ($\alpha = 0$). The corresponding phase diagram is richer, since superconductivity appears in addition to trivial magnetic orders. Because the spin-flip and pair-hopping terms in Eq. (2) are dropped in the spin-anisotropic case with $\alpha = 0$, we do not have to consider $\chi^{b,c}$ and the calculation of the two-particle Green functions simplifies substantially. The independent order parameters are also changed from the previous section, and are listed in Table II. We will consider only uniform ($q = 0$) and staggered ($q = Q$) ordering vectors.

TABLE II. Classification of independent operators for the Ising-anisotropic case ($\alpha = 0$) without CF splitting.

Identifier	Operator
(i')	n
(ii')	s^z
(iii')	s^x, s^y, o^{zx}, o^{zy}
(iv')	τ^z
(v')	$\tau^x, \tau^y, o^{xz}, o^{yz}$
(vi')	o^{zz}
(vii')	$o^{xx}, o^{xy}, o^{yx}, o^{yy}$
(viii')	p^{sx}, p^{sy}
(ix')	p^{zs}, p^{sz}
(x')	p^{xs}, p^{ys}

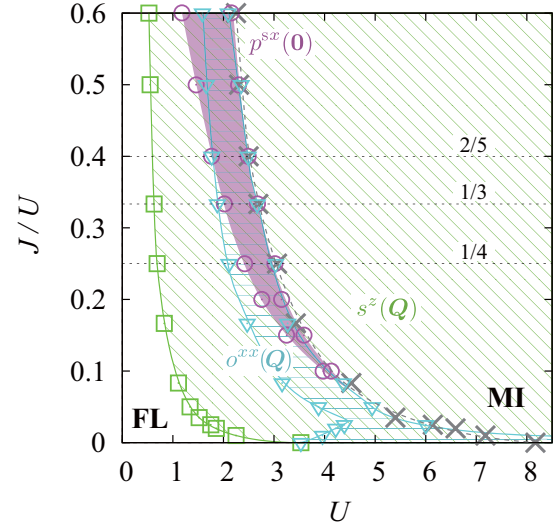


FIG. 2. Phase boundaries determined by the divergent points of susceptibilities at $n = 2$ and $T = 0.02$. The cross symbols indicate Mott transition points. The line $J/U = 1/4$ is the ratio considered in this paper, $J/U = 1/3$ marks the border from repulsive to attractive interactions, and $J/U = 2/5$ is a high-symmetry line (see text). The bold letters show Fermi liquid (FL) and Mott insulator (MI) phases without long-range orders.

In the remainder of this section, we focus on s^z , o^{xx} , and p^{sx} , since the corresponding symmetry breaking appears in regions which are not dominated by other ordering, for the parameters considered here. Figure 2 shows the phase diagram in the plane of U and J/U at half filling ($n = 2$) and $T = 0.02$. For $J = 0$, because of the high symmetry SU(4), all the diagonal orders except for the charge order are degenerate. On the other hand, a finite J substantially stabilizes the magnetic order with $s^z(Q)$. At the same time, the critical value of the metal-insulator transition is reduced by the Hund coupling. There appears staggered spin-orbital order [$o^{xx}(Q)$] and orbital-singlet/spin-triplet pairing [$p^{sx}(0)$], although these ordered regions are covered by the $s^z(Q)$ order. The spin-orbital order realized at $J = 0$ is destabilized by a small Hund coupling J . This is because the $c_{1\uparrow}^\dagger c_{2\downarrow}$ operator is relevant for the spin-orbital order with o^{xx} , and this spin-orbital flipping process is suppressed by a positive J which favors equal-spin states. However, it is stabilized near the Mott transition due to local spin and charge fluctuations. As will be shown later, these orders become most stable away from half filling or in the presence of a CF splitting.

It is notable that at $J/U = 0.4$ the orders with $o^{xx}(Q)$ and $p^{sx}(0)$ have the same transition point. This can be understood by symmetry considerations. We first define the particle-hole transformation for orbital 2,

$$\mathcal{P} c_{i2\sigma} \mathcal{P}^{-1} = \sum_{\sigma'} \sigma_{\sigma\sigma'}^\dagger c_{i2\sigma'}^\dagger e^{i\mathbf{Q} \cdot \mathbf{R}_i}, \quad (29)$$

which leaves $c_{i1\sigma}$ unchanged. This is similar to the transformation from repulsive to attractive interactions in the half-filled single-band Hubbard model [29]. One can show that this transformation does not change the Hamiltonian with Ising anisotropy ($\alpha = 0$) at $J/U = 2/5$ and $n = 2$. (In fact, this

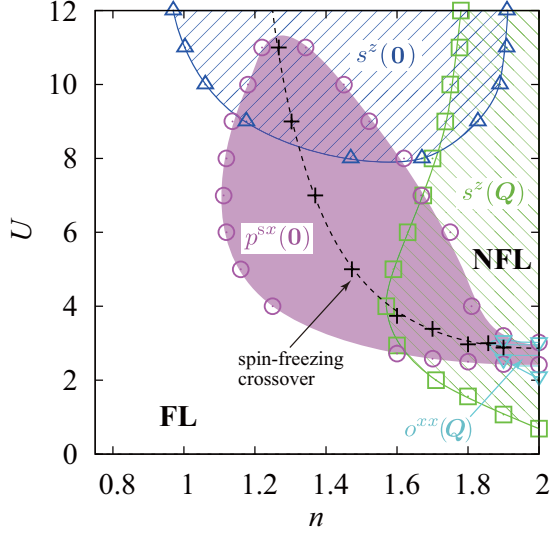


FIG. 3. Filling (n) vs interaction (U) phase diagram of the two-orbital model with $J/U = 1/4$ at $T = 0.02$. The Fermi-liquid (FL) and non-Fermi-liquid (NFL) regimes are separated by the spin-freezing crossover line.

invariance holds also for the case where the spin-flip term is included, but not in the presence of the pair-hopping term.) On the other hand, \mathcal{P} transforms the off-diagonal order into a diagonal order:

$$\mathcal{P} p^{sx}(\mathbf{0}) \mathcal{P}^{-1} = o^{xx}(\mathbf{Q}). \quad (30)$$

Thus, these two orders are degenerate at this particular value of J/U .

Figure 3 plots the phase diagram of the orbitally degenerate model away from half filling, for $T = 0.02$ and $J/U = 1/4$. Near half filling, an antiferromagnetically ordered phase is found, while in the large U regime, a ferromagnetic phase extends over a wide filling range. Between these two magnetically ordered regions, we find an instability toward an orbital-singlet spin-triplet superconducting state with $p^{sx}(\mathbf{0})$. Here, the quasilocal Cooper pair results from purely repulsive interactions, by a mechanism that has been proposed in the cold atom context [30]. This phase extends from the end-point of the half-filled Mott insulator along the so-called spin freezing line [10] into the metallic regime, in analogy to the three orbital results discussed in Ref. [4]. The spin-freezing phenomenon is well characterized by the local quantity

$$\Delta\chi_{\text{loc}} = \int_0^\beta d\tau [\langle s_i^z(\tau) s_i^z \rangle - \langle s_i^z(\beta/2) s_i^z \rangle]. \quad (31)$$

The first term is the local spin susceptibility. The long-time spin correlator, which appears as the second term, reflects the presence of frozen moments. The difference $\Delta\chi_{\text{loc}}$ thus quantifies the local spin fluctuations, whose maxima in parameter space define the spin-freezing crossover line [4]. This line is closely related to the spin-triplet superconductivity as shown in Fig. 3. While the main topic of this paper is the effect of CF splitting, Figs. 2 and 3 are important for a deeper understanding of the resulting ordered states under the CF, as discussed in the following.

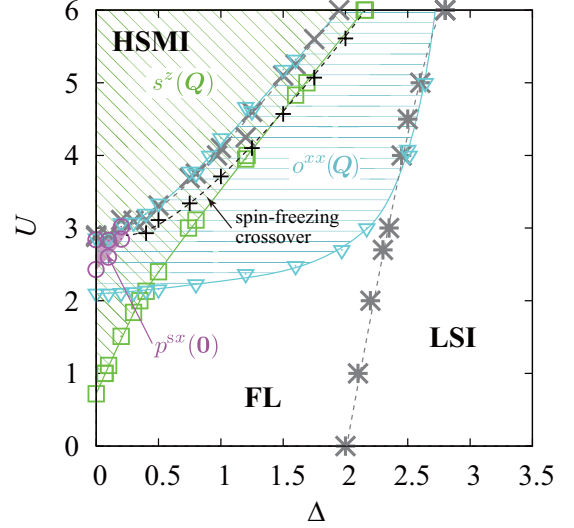


FIG. 4. Phase diagram of the half-filled two-orbital model with $J/U = 1/4$ in the space of CF splitting Δ and U at temperature $T = 0.02$. In the disordered case, we have Fermi-liquid (FL), high-spin Mott insulator (HSMI), and low-spin insulator (LSI) phases.

B. Split orbitals

We next consider the half-filled two-orbital model with an additional crystal field (CF) splitting given by

$$\mathcal{H}_{\text{CF}} = \Delta \sum_{i\sigma} (n_{i1\sigma} - n_{i2\sigma}). \quad (32)$$

For $\alpha = 1$, the metal-insulator phase diagram of this model without ordering has been studied in Ref. [5]. A qualitatively similar diagram is obtained also for the $\alpha = 0$ case [31]. At large U and small Δ , there is a high-spin (Mott) insulating phase with small orbital polarization. For sufficiently large Δ , the system is in a low-spin insulating state with large orbital polarization, which for $U \rightarrow 0$ is adiabatically connected to the band insulator. At large enough U and finite T , there is a transition from the high-spin to the low-spin insulator at $\Delta \simeq 3J/2$ [11]. At weaker U , there exists an intermediate metallic phase, which for large J/U and low T extends along the $\Delta \simeq 3J/2$ line to rather large values of Δ .

Figure 4 shows the Δ - U phase diagram for the anisotropic case ($\alpha = 0$) at half filling. While the high-spin insulator is antiferromagnetically ordered, the metallic nose separating the high-spin and low-spin insulators is unstable against the staggered spin-orbital ordering with $o^{xx}(\mathbf{Q})$. This result is consistent with the excitonic insulator state found in this parameter regime in Ref. [8]. The low-spin insulating region is not susceptible to long-range order. We also find an instability toward the staggered high-spin/low-spin order reported in Ref. [7]. This instability, which can be detected by the orbital susceptibility with $\tau^z(\mathbf{Q})$, appears for $U \gtrsim 6$. However, we do not show this phase, since it is covered by the $o^{xx}(\mathbf{Q})$ order at least for $U \leq 8$.

We now discuss the connections between the two phase diagrams in Figs. 3 and 4, which follow from the symmetry relation (30). Away from half filling, in the model without CF splitting, the spin-triplet superconductivity $p^{sx}(\mathbf{0})$ is stabilized

along the spin-freezing crossover line, which indicates that local spin fluctuations induce the pairing state [4]. If we perform the particle-hole transformation, this pairing state changes into the ordered state with $o^{xx}(\mathbf{Q})$ as discussed above. At the same time, the chemical potential term $\mathcal{H}_{\text{chem}} = -\mu \sum_{i\gamma\sigma} n_{i\gamma\sigma}$ is transformed into a CF splitting term:

$$\mathcal{P} \mathcal{H}_{\text{chem}} \mathcal{P}^{-1} = -\mathcal{H}_{\text{CF}} \quad (33)$$

with $\Delta = \mu$. In other words, *the spin-triplet superconductivity away from half filling corresponds to the spin-orbital order under the CF splitting*. Furthermore the local spin moment is unchanged: $\mathcal{P} s_i^z \mathcal{P}^{-1} = s_i^z$. Hence, in both cases the local spin fluctuations play an important role in stabilizing the ordered state. As shown in Fig. 4, there is indeed a spin-freezing line extending along the metallic nose in the region where spin-orbital order appears. While this mapping is exact only for $J/U = 2/5$, the qualitative correspondence between the two physical situations should be valid in a broader regime, including the case $J/U = 1/4$ considered in Figs. 3 and 4.

V. THREE-ORBITAL MODEL

A. General remarks

We are interested in the stability of the interorbital-spin-triplet ($J > 0$) and intraorbital-spin-singlet ($J < 0$) superconducting phases to CF splittings of the “2/1” type,

$$\mathcal{H}_{\text{CF}} = -\Delta \sum_{i\sigma} (n_{i1\sigma} + n_{i2\sigma} - n_{i3\sigma}). \quad (34)$$

This term breaks the orbital $\text{SO}(3)$ symmetry, but the $\text{SO}(2)$ symmetry within the $\gamma = 1, 2$ subspace remains for $\alpha = 1$.

At half filling and without CF splitting, the Hamiltonian is invariant under the simple particle-hole transformation defined by

$$\mathcal{P}_0 c_{i\gamma\sigma} \mathcal{P}_0^{-1} = c_{i\gamma\sigma}^\dagger e^{i\mathbf{Q} \cdot \mathbf{R}_i}, \quad (35)$$

i.e., we have $\mathcal{P}_0 \mathcal{H} \mathcal{P}_0^{-1} = \mathcal{H}$. On the other hand, the CF Hamiltonian is transformed as $\mathcal{P}_0 \mathcal{H}_{\text{CF}} \mathcal{P}_0^{-1} = -\mathcal{H}_{\text{CF}}$. Thus the sign of the CF splitting Δ is reversed. For some interaction and filling parameters, as seen below, the present system shows superconductivity with interorbital spin-triplet pairs,

$$p_t^{\gamma\gamma'}(\mathbf{0}) = \sum_i c_{i\gamma\uparrow}^\dagger c_{i\gamma'\uparrow} + \text{H.c.}, \quad (36)$$

for $J > 0$ and $\gamma \neq \gamma'$, or intraorbital spin-singlet pairs,

$$p_s^\gamma(\mathbf{0}) = \sum_i c_{i\gamma\uparrow}^\dagger c_{i\gamma\downarrow} + \text{H.c.}, \quad (37)$$

for $J < 0$. Here we choose these expressions for the pair amplitudes, although we can classify them using a similar method as given in Eqs. (11) and (12) using Pauli and Gell-Mann matrices. We have the symmetry relations $\mathcal{P}_0 p_t^{\gamma\gamma'} \mathcal{P}_0^{-1} = -p_t^{\gamma\gamma'}$ and $\mathcal{P}_0 p_s^\gamma \mathcal{P}_0^{-1} = -p_s^\gamma$. While the signs are reversed after the transformation, i.e., the phases of the pair amplitudes are rotated by π , their forms are unchanged after the particle-hole transformation.

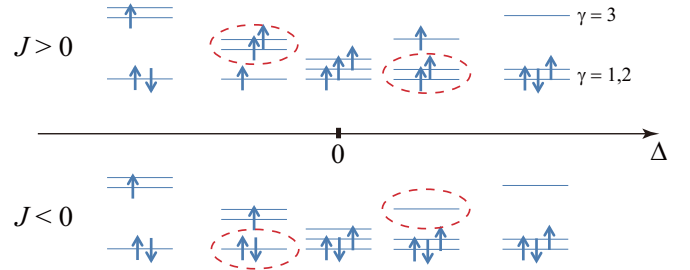


FIG. 5. Schematic illustration of atomic configurations as a function of Δ at half filling ($n = 3$). The circles with dotted lines show the orbital which is responsible for superconductivity.

To understand the implications of the above symmetry relations, let us first consider the $J > 0$ model at half filling. In this case the same-spin electrons tend to occupy the same site to form Cooper pairs. At $\Delta = 0$, the three pairs p_t^{12} , p_t^{23} , and p_t^{31} are degenerate. For $\Delta > 0$, as illustrated in the top part of Fig. 5, one expects that p_t^{12} is more stable than p_t^{23} and p_t^{31} because the orbitals with $\gamma = 1, 2$ have a lower energy than $\gamma = 3$. In a similar manner one may also speculate that for $\Delta < 0$ the pairs p_t^{23} and p_t^{31} are more stable than p_t^{12} . However this is *incorrect*. The pair p_t^{12} is the most stable even for $\Delta < 0$. This is because the pairing amplitudes are the same for $\Delta > 0$ and $\Delta < 0$, according to the above particle-hole symmetry argument, even though the Cooper pairs are formed in the higher-energy orbital (broken circle in the upper panel of Fig. 5) when $\Delta < 0$. This at first sight counterintuitive result can be rationalized by considering the degeneracy: the p_t^{23} and p_t^{31} pairs are destabilized by fluctuations among these energetically degenerate states, which may diminish the pairing compared to the nondegenerate case. On the other hand, the p_t^{12} pairs are not subject to such fluctuations and are therefore more stable.

One can also think of other effects which help explain this peculiar behavior. Whereas the lower-energy orbital is easily occupied and pairs between low-energy and high-energy electrons can be formed for $\Delta < 0$, the mobility of these pairs, which is also important for realizing superconductivity, is reduced because of the high occupancy of the low-energy orbitals. On the other hand, for higher-energy orbitals the mobility of the pairs is high, although the pairs themselves are harder to form. As a result, the superconducting state resulting from pairs in the high-energy orbitals is actually more robust, which is also supported by the numerical results as shown later.

The same argument can also be applied to the case of $J < 0$. At $\Delta = 0$, the three pairs p_s^1 , p_s^2 , and p_s^3 are degenerate. For $\Delta < 0$, as illustrated in the lower panel of Fig. 5, one expects that p_s^3 is more stable than p_s^1 and p_s^2 . On the other hand, for $\Delta > 0$ one might speculate that p_s^1 and p_s^2 are more stable than p_s^3 , but this is again incorrect. Although the level of the orbital $\gamma = 3$ is higher than the others, the most stable Cooper pair is formed in this high-energy orbital.

Note that in the above discussion, we do not invoke any subtleties of the system, only the particle-hole symmetry of the original Hamiltonian without CF splitting. Keeping these facts in mind, we can readily understand the result in the following subsections.

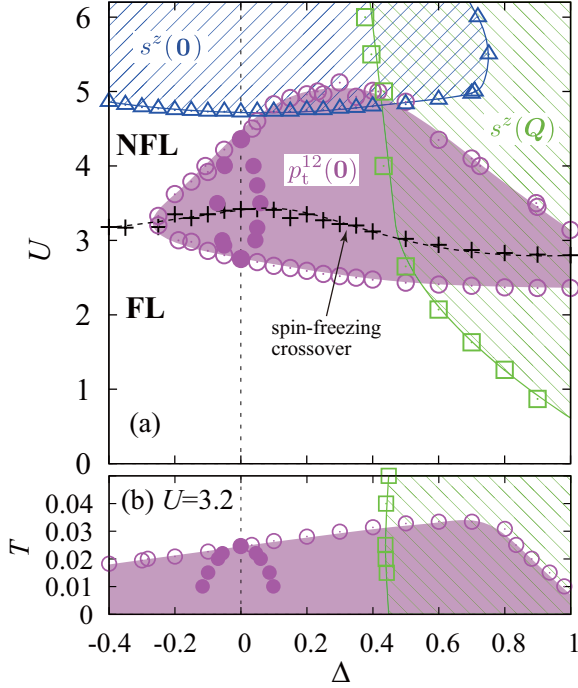


FIG. 6. (a) Phase boundaries determined by the divergence of susceptibilities for the parameters $n = 2$, $J/U = 1/4$, and $T = 0.02$. Open (filled) circles correspond to phase boundaries for $p_t^{12}(p_t^{23,31})$ pairing. The limit $\Delta \rightarrow +\infty$ corresponds to the two-orbital model at half filling. (b) Phase diagram in the temperature (T)-CF splitting (Δ) space for $U = 3.2$.

B. $J > 0$ case

In the remainder of this section, we show results for the three-orbital model with CF splitting, focusing again on the case with Ising anisotropy ($\alpha = 0$). For $J > 0$, the pairing amplitude is given by Eq. (36), and the magnetic moments for the three-orbital model are defined by

$$s^z(\mathbf{q}) = \sum_{i\gamma\sigma} c_{i\gamma\sigma}^\dagger \sigma_{\sigma\sigma}^z c_{i\gamma\sigma} e^{-i\mathbf{q}\cdot\mathbf{R}_i}. \quad (38)$$

Other ordered states are not considered here since these are covered by the orders $p_t^{\gamma\gamma'}(\mathbf{0})$ and $s^z(\mathbf{q})$ in the parameter range considered in this paper.

At $\Delta = 0$ the phase diagram for the three-orbital model [4] has properties similar to those of the two-orbital model shown in Fig. 3. Here we discuss the effect of the CF splitting on the three-orbital model with $n = 2$, because it is relevant to Sr_2RuO_4 as discussed later. Figure 6(a) shows the U - Δ phase diagram at $n = 2$ and $T = 0.02$. Note that we do not have a symmetry between $\Delta > 0$ and $\Delta < 0$ for the case away from half filling. The antiferromagnetic phase [$s^z(\mathbf{Q})$] dominates for sufficiently large Δ because in the limit $\Delta \rightarrow \infty$ the system becomes the half-filled two-orbital model discussed in the previous section. In the large- U region the ferromagnetic order [$s^z(\mathbf{0})$] appears.

The interorbital spin-triplet superconductivity is realized at intermediate values of U and for a large range of Δ . The spin-freezing crossover defined by the maximum of Eq. (31) is also plotted, and it is obvious that the superconductivity is

stabilized near this line. For $\Delta \neq 0$, the degeneracy of the three pairs is lifted, and the most stable one is p_t^{12} regardless of the sign of Δ , as explained in the previous subsection. The other pairings $p_t^{23,31}$ (filled circles in Fig. 6) are quickly suppressed.

The CF splitting dependence of the transition temperature is shown in Fig. 6(b). It is notable that the transition temperature is enhanced by the CF splitting for $\Delta > 0$. This behavior might be due to a reduction of fluctuations among orbitals and an enhanced probability of pair formation by the splitting of the degenerate orbitals. For $\Delta < 0$ such an enhancement is not seen, presumably because the most stable Cooper pair is in this case formed in the high-energy orbitals; this pair formation is hence destabilized by the CF splitting. Still, the p_t^{12} interorbital spin-triplet superconductivity can be observed in a wide region of $|\Delta|$ at low temperatures.

Our simulations are relevant for strontium ruthenate compounds. In Sr_2RuO_4 , the filling is $n = 4$, which is identical to $n = 2$ due to particle-hole symmetry, and the interaction is estimated as $U \simeq 3.2$ [32]. In this compound the CF splitting among the three t_{2g} orbitals has the form of Eq. (34) due to the tetragonal symmetry of the crystal. According to Figs. 6, the spin-triplet superconductivity occurs at low temperatures even for finite CF splittings in the physically reasonable range. The pairing is most robust for same-spin electrons in the degenerate orbitals. We thus obtain consistent results for this strontium ruthenate compound, although for a more detailed comparison with experiments we should consider the realistic band structure.

Here we briefly comment on the effect of α which restores the isotropy. For the present spin-triplet superconductivity with $J > 0$, the effect of spin-flip terms is relevant and suppresses this pairing due to the fluctuations among the three spin-triplet states [4]. Indeed we could not find the spin-triplet superconductivity down to the lowest accessible temperatures for $\alpha = 1$. Thus the anisotropy in spin space is important for its realization. Despite this fact, the spin-triplet superconducting state can be realized in Sr_2RuO_4 , since only a small anisotropy is necessary for its stabilization [4].

C. $J < 0$ case

Because of the relevance for fulleride compounds, it is also interesting to consider the half-filled ($n = 3$) three-orbital model with negative Hund coupling [24]. In addition to the magnetic moment defined in the last subsection, we consider two kinds of orbital moments defined by

$$\tau^{3,8}(\mathbf{q}) = \sum_{i\gamma\sigma} c_{i\gamma\sigma}^\dagger \lambda_{\gamma\gamma}^{3,8} c_{i\gamma\sigma} e^{-i\mathbf{q}\cdot\mathbf{R}_i}, \quad (39)$$

where $\lambda^3 = \text{diag}(1, -1, 0)$ and $\lambda^8 = \text{diag}(1, 1, -2)/\sqrt{3}$ are Gell-Mann matrices. These orbital orders are degenerate for $\Delta = 0$, but become different in the presence of CF splitting. For the parameter range considered in this paper, the orbital order with $\tau^3(\mathbf{Q})$ is more stable than the other, so we plot only the phase boundaries for $\tau^3(\mathbf{Q})$.

Figure 7(a) shows the U - Δ phase diagram at $J/U = -1/10$ and $T = 0.02$. Although this ratio of J/U is much larger than the estimate for alkali-metal-doped fullerenes [23], it is suitable for clarifying the qualitative behaviors originating from the negative Hund coupling. Without any long-range

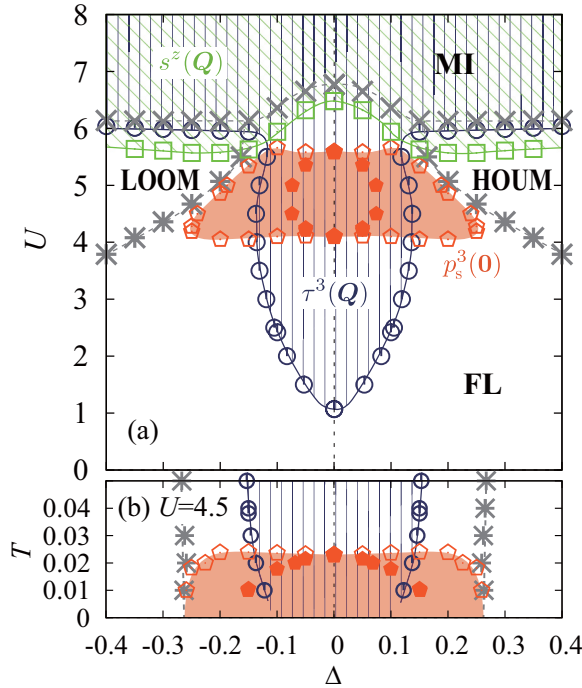


FIG. 7. (a) Phase boundaries determined by the divergence of susceptibilities for $n = 3$, $J/U = -1/10$, and $T = 0.02$. The $\Delta < 0$ part is equivalent to the $\Delta > 0$ part due to the particle-hole symmetry at half filling. Here, in addition to Fermi liquid (FL) and Mott insulator (MI) phases, we have a higher-orbital unoccupied metal (HOUM) for $\Delta > 0$ and a lower-orbital occupied metal (LOOM) for $\Delta < 0$. Open (filled) pentagons correspond to p_s^3 ($p_s^{1,2}$) pairing. (b) Phase diagram in the CF splitting (Δ)-temperature (T) space for $U = 4.5$.

order, we have three kinds of states depending on the parameters. In the small and large U regions, metallic and Mott insulating states are realized, respectively. For sufficiently large CF splitting, a new state appears in between, which is a metallic state with a completely empty (occupied) orbital for $\Delta > 0$ ($\Delta < 0$) [33].

Now we discuss the ordered states. The orbital degrees of freedom are activated by the negative Hund coupling and are ordered in a staggered manner as shown in Fig. 7(a). Orbital order with $\tau^3(Q)$ moments is the dominant phase near $\Delta = 0$. For large U , antiferromagnetism is also stabilized, which is explained by the still remaining active spin degrees of freedom in the half-filled system.

As shown in Fig. 7, the intraorbital spin-singlet pairing with p_s^3 is realized in the intermediate U region. This pairing is also caused by the negative Hund coupling, which favors doubly occupied orbitals. The other pairings with $p_s^{1,2}$ are less stable (see filled pentagons in the figure) as discussed before. For small Δ , this superconducting state is covered by the orbital order. With increasing CF splitting, we find a region between the orbital ordered phase and the occupied/unoccupied orbital metal, where the p_s^3 pairing state is most stable. The transition temperature as a function of Δ is plotted in Fig. 7(b). It is notable that the transition temperature is slightly enhanced by the CF splitting. This enhancement can be intuitively interpreted again as resulting from the suppression of fluctuations among orbitals and an increasing probability of pair formation for

$\Delta < 0$. At the same time, the mobility of the pairs is reduced, so that T_c is almost independent of Δ . For $\Delta > 0$, on the other hand, the probability of pair formation instead of the mobility of pairs is decreased, to give the exactly same result as in the $\Delta < 0$ case. The p_s^3 pairing state disappears when orbital $\gamma = 3$ becomes fully occupied or empty.

Finally we discuss the effect of the parameter α . In the $J < 0$ case, the pair hopping is relevant and the spin flip is not. According to Ref. [24], the pair hopping substantially enhances the intraorbital spin-singlet pairing. Hence the transition temperature should increase for $\alpha > 0$, which is in contrast to the $J > 0$ case. While the realistic value of J is tiny ($J/U \sim -0.025$ for fullerenes [23]), the pairing from negative Hund coupling has this advantage and has a chance to be realized in systems with isotropic interaction ($\alpha = 1$). The effect of the increasing orbital fluctuations on the intraorbital spin-singlet superconducting state for $\alpha > 0$, however, remains to be investigated.

VI. SUMMARY AND DISCUSSION

We have presented a systematic analysis of the electronic ordering instabilities in two- and three-orbital Hubbard models. We have used DMFT in combination with a numerically exact hybridization expansion impurity solver and a semicircular density of states. Our results thus represent the generic phase diagrams of high-dimensional lattice models, irrespective of details of the band structure. Only uniform and staggered order parameters have been considered, and we focused on the intermediate coupling regime, which is relevant for most unconventional multiband superconductors and cannot be accessed reliably by approximate methods such as the fluctuation-exchange approximation.

In the case with positive Hund coupling, we found instabilities to antiferromagnetic and ferromagnetic orders, spin-orbital order, and orbital-singlet spin-triplet superconductivity. In the model with negative Hund coupling we identified antiferro-orbital order, antiferromagnetism, and intraorbital spin-singlet pairing. We have shown how symmetry relations can be used to connect some of these ordered phases, and to understand the Cooper pairings in three-orbital models with CF splittings.

In particular, we showed that in the two-orbital model with $J/U = 2/5$, the orbitally degenerate system away from half filling can be mapped exactly onto the half-filled system with CF splitting, and that the orbital-singlet spin-triplet superconductivity in the former system corresponds to spin-orbital order in the latter. The qualitative correspondence between these at first sight different physical situations can be expected to hold in a wider parameter regime. Since the fluctuating local moments at the border of the spin-frozen metal regime play a crucial role in stabilizing the spin-triplet superconducting state, and the spin moments are unaffected by the mapping, this symmetry argument also implies that fluctuating moments drive the instability to spin-orbital order. Thus the diagonal and off-diagonal orders can be understood in a unified manner.

Crystal-field splittings of the “2/1” type in the three orbital model can have a surprising effect on the superconducting state. In the spin-triplet state with filling $n = 2$ and $J > 0$, the most stable pairs are always formed in the two degenerate

orbitals, even if the energy of these orbitals is increased by the CF. On the other hand, in the half-filled spin-singlet pairing state with $J < 0$, the most stable pairs are always formed in the nondegenerate orbital, irrespective of the sign of the CF. We explained this behavior using a particle-hole transformation, and also gave an intuitive interpretation based on fluctuations of the condensate and the mobility of Cooper pairs.

The use of symmetry relations is also interesting from a computational point of view, since the rotational invariance of the Slater-Kanamori interaction allows us to express certain four-point correlation functions in terms of others which can be easily measured in standard hybridization expansion Monte Carlo simulations [15]. This trick however fails as soon as the rotational invariance is broken, e.g., by the presence of CF splittings. Whether or not it is possible to extend this technique to more general situations by exploiting additional symmetries is an interesting open problem. An alternative strategy is to directly measure all types of four-point correlation functions using a worm sampling algorithm [34].

Our three-orbital calculations with filling $n = 2$ (equivalent to $n = 4$) can be regarded as toy model simulations of Sr_2RuO_4 . This material is a quasi-2D system with tetragonal symmetry and has a “2/1” type CF splitting. While the CF splitting quickly suppresses the superconductivity in the nondegenerate orbital, the pairing between the degenerate orbitals is robust. Our calculations are consistent with a fluctuating-moment induced spin-triplet superconductivity in this compound, which has a partial orbital degeneracy (zx and yz orbitals). More accurate simulations would need to take into account the realistic band structure.

The half-filled three-orbital model with negative Hund coupling is of interest in connection with fulleride compounds, where the Jahn-Teller screening of the small J leads to a stabilization of the low-spin states [12]. The effect of CF splitting is to further stabilize the spin-singlet pairing in the nondegenerate orbital, and to suppress a competing orbital ordered phase. While the J parameters used in our study are larger than the *ab initio* estimates [23], this finding has some implications on the recently reported light-induced superconductivity in K_3C_{60} [25]. One possible explanation for the enhanced T_c , proposed in Ref. [25], is that the driving of a phonon of T_{1u} symmetry leads to an essentially static distortion of the C_{60} molecules, and hence a splitting of the initially degenerate molecular orbitals. Our result shows that this splitting, if studied within an equilibrium formalism, can contribute to a slightly enhanced pairing. The related effect of orbital differentiation in the interaction parameters still needs to be systematically explored. Given the large enhancement of T_c observed in the experiments, one may speculate that the J parameter is effectively modified in the driven state (enhanced dynamical Jahn-Teller effect). This, and other nonequilibrium phenomena should be investigated within a Floquet formalism or using the nonequilibrium extension of DMFT [35].

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APPENDIX A: BETHE LATTICE IN INFINITE DIMENSIONS

Here we consider some properties of the Bethe lattice in infinite dimension. Since the wave vectors are ill defined in this special lattice, we have to deal with real space. However, we will show that the concept of wave vectors can be partially applied, and that a treatment similar to that of ordinary lattices is possible.

1. Single-particle Green function

Let us here consider the tight-binding model for noninteracting spinless fermions on the Bethe lattice:

$$\mathcal{H}_0 = -t \sum_{(ij)} c_i^\dagger c_j, \quad (\text{A1})$$

where the summation with respect to the site index is over the nearest-neighbor pairs. We begin with the equations of motion given by

$$-\partial_\tau c_i = -t \sum_\delta c_{i+\delta}, \quad (\text{A2})$$

$$-\partial_\tau c_{i+\delta} = -t c_i - t \sum_{\delta+\delta' \neq 0} c_{i+\delta+\delta'}, \quad (\text{A3})$$

where $-\partial_\tau O = [O, \mathcal{H}_0]$. The site index $i + \delta$ denotes the nearest-neighbor sites of the site i . The Fourier transformation of the equations of motion for the Green function defined by $G_{ij}(\tau) = -\langle T_\tau c_i(\tau) c_j^\dagger \rangle$ is given by

$$z G_0(z) = 1 - t d G_1(z), \quad (\text{A4})$$

$$z G_k(z) = -t G_{k-1}(z) - t d G_{k+1}(z), \quad (\text{A5})$$

for $k = 1, 2, \dots$, which represents the number of sites between the positions at i and j . Here z is a complex frequency, d is a connectivity, and G_0 is a local Green function. We have taken the limit $d \rightarrow \infty$. Defining $\alpha_k = G_k/G_{k-1}$, one can show the following relation:

$$\alpha_k = \frac{-t}{z + t d \alpha_{k+1}} = \frac{-t}{z + t d \frac{-t}{z + t d \frac{-t}{z + \dots}}} = \frac{-t}{z + t d \alpha_k}. \quad (\text{A6})$$

Hence the ratio α_k is independent of k , and we can write it as $\alpha_k = \alpha$. This quantity is explicitly derived by solving the quadratic equation and we obtain the Green function as

$$G_k(z) = -\alpha(z)^{k+1}/t. \quad (\text{A7})$$

One of the two solutions for α is chosen so that it behaves as $G_0(z) \rightarrow 1/z$ when $|z| \rightarrow \infty$. The form of the Green function is the same as the one obtained in Ref. [36]. For interacting systems, the local self-energy Σ is included by the replacement $z \rightarrow z - \Sigma$.

2. Two-particle Green function

We define the two-particle Green functions without vertex parts by

$$\chi_{\text{unif}}^0(z) = -\frac{1}{N} \sum_{ij} G_{ij}(z) G_{ji}(z), \quad (\text{A8})$$

$$\chi_{\text{stag}}^0(z) = -\frac{1}{N} \sum_{ij} \lambda_i \lambda_j G_{ij}(z) G_{ji}(z), \quad (\text{A9})$$

which are relevant to uniform and staggered susceptibilities, respectively. Here $\lambda_i = \pm 1$ is a sign which depends on the sublattice. Substituting the results in the previous subsection, we obtain

$$\chi_{\text{unif}}^0 = -\frac{(\alpha/t)^2}{1 - d\alpha^2}, \quad \chi_{\text{stag}}^0 = -\frac{(\alpha/t)^2}{1 + d\alpha^2}. \quad (\text{A10})$$

This leads to the simpler expressions

$$\chi_{\text{unif}}^0 = \frac{dG_0}{dz}, \quad \chi_{\text{stag}}^0 = -\frac{G_0}{z}. \quad (\text{A11})$$

Thus we derive the uniform and staggered components of two-particle Green functions.

We now consider another representation of the above two-particle Green functions, and introduce a density of states and a “wave-vector \mathbf{k} summation” by

$$G_0(z) = \int \frac{\rho(\varepsilon)}{z - \varepsilon} d\varepsilon \equiv \frac{1}{N} \sum_{\mathbf{k}} g_{\mathbf{k}}(z), \quad (\text{A12})$$

where $\rho(-\varepsilon) = \rho(\varepsilon)$, $g_{\mathbf{k}}(z) = 1/(z - \varepsilon_{\mathbf{k}})$, and $N = \sum_{\mathbf{k}} 1$. Here we do not have to know the specific form of $\varepsilon_{\mathbf{k}}$, and only assume the existence of a vector \mathbf{Q} defined by $\varepsilon_{\mathbf{k}+\mathbf{Q}} = -\varepsilon_{\mathbf{k}}$. We also introduce the \mathbf{q} -dependent two-particle Green function by

$$\tilde{\chi}_{\mathbf{q}}^0(z) = -\frac{1}{N} \sum_{\mathbf{k}} g_{\mathbf{k}}(z) g_{\mathbf{k}+\mathbf{q}}(z). \quad (\text{A13})$$

Using Eq. (A12), one can show the relations $\tilde{\chi}_{\mathbf{0}}^0 = \chi_{\text{unif}}^0$ and $\tilde{\chi}_{\mathbf{Q}}^0 = \chi_{\text{stag}}^0$. Thus the pseudo-wave-vectors $\mathbf{q} = \mathbf{0}$ and

$\mathbf{q} = \mathbf{Q}$ correspond to the uniform and staggered components, respectively. Namely, the Bethe lattice can be partly handled as if it were a simple cubic lattice which has the same relation $\varepsilon_{\mathbf{k}+\mathbf{Q}} = -\varepsilon_{\mathbf{k}}$ with the staggered ordering vector $\mathbf{Q} = (\pi, \pi, \pi)$. Note that this analogy is valid only for the uniform and staggered components.

APPENDIX B: PHYSICAL INTERPRETATION OF ORDER PARAMETERS

Let us physically interpret the orbital and spin-orbital moments in terms of doubly degenerate e_g orbitals for d electrons under the cubic symmetry. The wave functions are written as

$$|\gamma = 1\rangle = |0\rangle, \quad (\text{B1})$$

$$|\gamma = 2\rangle = (|2\rangle + |-2\rangle)/\sqrt{2}, \quad (\text{B2})$$

where $|m\rangle$ on the right-hand side is the eigenstate of the angular momentum ℓ_z . With use of this expression, we can show the relations

$$\tau^z \propto 3\ell_z^2 - \ell^2, \quad (\text{B3})$$

$$\tau^x \propto \ell_x^2 - \ell_y^2, \quad (\text{B4})$$

$$\tau^y \propto \overline{\ell_x \ell_y \ell_z}, \quad (\text{B5})$$

where the overline symmetrizes the product of operators as, e.g., $\ell_x \ell_y = (\ell_x \ell_y + \ell_y \ell_x)/2$. This can be shown by standard quantum-mechanics calculations for angular momentum. Thus τ^z and τ^x are rank-2 operators, while τ^y is a rank-3 operator. For the spin-orbital moment, we further put the spin moment on the orbital moment τ^μ , and hence the rank of the operator is increased by 1. Namely, $o^{z\mu}$ and $o^{x\mu}$ are rank-3 operators, and $o^{y\mu}$ is a rank-4 operator. In the context of f -electron systems, rank-0, -1, -2, -3, -4 operators are called “scalar (monopole),” “dipole,” “quadrupole,” “octupole,” and “hexadecapole,” respectively [37]. The odd (even) rank tensor is time-reversal odd (even).

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