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Novel Block Excitonic Condensate at $n = 3.5$ in a Spin-Orbit Coupled t_{2g} Multiorbital Hubbard Model

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Theoretical studies recently predicted the condensation of spin-orbit excitons at momentum $q=\pi$ in t_{2g}^4 spin-orbit coupled three-orbital Hubbard models at electronic density $n = 4$. In parallel, experiments involving iridates with non-integer valence states for the Ir ions are starting to attract considerable attention. In this publication, using the density matrix renormalization group technique we present evidence for the existence of a novel excitonic condensate at $n = 3.5$ in a one-dimensional Hubbard model with a degenerate t_{2g} sector, when in the presence of spin-orbit coupling. At intermediate Hubbard U and spin-orbit λ couplings, we found an excitonic condensate at the unexpected momentum $q=\pi/2$ involving $j_{\text{eff}} = 3/2, m = \pm 1/2$ and $j_{\text{eff}} = 1/2, m = \pm 1/2$ bands in the triplet channel, coexisting with an also unexpected block magnetic order. We also present the entire λ vs U phase diagram, at a fixed and robust Hund coupling. Interestingly, this new “block excitonic phase” is present even at large values of λ , unlike the $n = 4$ excitonic phase discussed before. Our computational study helps to understand and predict the possible magnetic phases of materials with $d^{3.5}$ valence and robust spin-orbit coupling.

I. INTRODUCTION

In the last decade, the $4d/5d$ transition metal oxides have received considerable attention in the Condensed Matter community, specially because they provide a unique platform for the development of unconventional magnetic and transport properties mainly as a consequence of a robust spin-orbit coupling [1–11]. One of the most interesting materials is Sr_2IrO_4 containing Ir^{4+} ions, with an electronic density $n = 5$ [12]. This compound displays similarities with La_2CuO_4 , even with a relatively smaller Hubbard repulsion, because both exhibit long-range antiferromagnetic ordering in quasi two-dimensional layers [13, 14]. The realization of an effective layered half-filled Hubbard model in Sr_2IrO_4 is a result of a spin-orbit coupling λ close to 0.5 eV that splits the t_{2g} states into $j_{\text{eff}} = 1/2$ and $j_{\text{eff}} = 3/2$ sectors with a gap approximately equal to $3\lambda/2$. At $n = 5$ this leads to a half-filled $j_{\text{eff}} = 1/2$ band and concomitant Mott/Slater insulator behavior. Besides the iridates, other materials have also similarly interesting properties [15–17]. Even in the context of iron superconductors the importance of spin-orbit coupling has been remarked [18–21].

Another interesting scenario which has been recently theoretically investigated led to the prediction of unusual magnetism in the $n = 4$ case [22–28]. At this electronic density, *spin-orbit excitons* (for details see Sec. III A) were found to condense at momentum $q = \pi$, both in the intermediate and strong coupling limits, and also display antiferromagnetic staggered magnetic order. Experimentally, for double perovskite materials such as Sr_2YIrO_6 and Ba_2YIrO_6 , with Ir^{5+} ions and a $5d^4$ configuration, the presence of the exciton condensate, as discussed time

ago in semiconductors [29], has been debated [30–34]. Recent RIXS (resonant inelastic x-ray scattering) experiments on Sr_2YIrO_6 and Ba_2YIrO_6 have unveiled $J_{\text{eff}} = 1$ and $J_{\text{eff}} = 2$ excitations with weak dispersion at energies approximately 0.37 eV and 0.7 eV [35], respectively, which suggests that the bandwidth of excitonic excitations is not sufficiently large when compared with λ to realize the predicted spin-orbit exciton condensate. It should be noted that these $J_{\text{eff}} = 1$ and $J_{\text{eff}} = 2$ excitations can be understood in terms of more conventional excitonic (electron-hole pair) states [29] between $j_{\text{eff}} = 3/2$ and $j_{\text{eff}} = 1/2$ sectors. Because in these excitations electrons jump from $j_{\text{eff}} = 3/2$ to $j_{\text{eff}} = 1/2$ states, the addition of angular momentum suggests that this will lead to $J_{\text{eff}} \in \{\frac{3}{2} - \frac{1}{2}, \frac{3}{2} + \frac{1}{2}\}$ i.e. $J_{\text{eff}} = 1$ or 2 excitations. In the layered Sr_2IrO_4 compound, these spin-orbit excitons are also present as stable excited states as shown by recent RIXS and optical conductivity measurements [36–38] [note that the notation J_{eff} is used for the total effective angular momentum of the system (or an atom), while j_{eff} refers for the effective angular momentum of single particle states. In the rest of the paper, we follow the same convention].

In addition to the above mentioned progress, it should be remarked that there are several real quasi-one dimensional materials with robust spin-orbit coupling strength that have been studied in the literature. The doped variants of the materials reported below may directly realize the physics discussed in this publication, because our calculations are based on numerically exact solutions of one-dimensional multiorbital models. For example, recently 1D stripes of Sr_2IrO_4 [39] were grown epitaxially and RIXS spectra have shown the presence of spin-

orbit excitons at energies nearly 0.6 eV. Other examples of one-dimensional $j_{\text{eff}} = 1/2$ antiferromagnets includes CaIrO_3 [40–43] and Ca_4IrO_6 [44, 45]. BaIrO_3 also belongs to the $5d^5$ class but have shown an unexpected charge-density wave [46, 47]. There are also mixed $3d$ - $5d$ one-dimensional insulators, such as $\text{Ba}_5\text{CuIr}_3\text{O}$ [48] and $\text{Sr}_3\text{CuIrO}_6$ [49, 50]. La_3OsO_7 , which lies in the category of $5d^3$, is also a quasi-one-dimensional material with antiferromagnetic ordering and $T_N = 45$ K [51]. There are also examples of quasi-one-dimensional materials with fractional valence states of the Ir and Rh ions, such as $\text{Ba}_5\text{AlIr}_2\text{O}_{11}$ [52–54], $\text{Ca}_5\text{Ir}_3\text{O}_{12}$ [55, 56], and $\text{Sr}_3\text{Rh}_4\text{O}_{12}$ [44]. $\text{BaRu}_6\text{O}_{12}$ and KRu_4O_8 are examples of quasi-one-dimensional ruthenates [57, 58] that have also attracted considerable attention. The combination of the existence of real quasi-one-dimensional $4d$ and $5d$ materials and our model studies employing numerically very accurate techniques provides a unique opportunity to explore and understand the phases which can emerge from the interplay of spin-orbit coupling, Coulomb electronic repulsion, and kinetic energy.

To obtain our results we use the numerically accurate density matrix renormalization group (DMRG) technique [59, 60] to solve the degenerate three-orbital Hubbard model in one-dimension. Up to now studies of the phases emerging in fractionally-filled three-orbital Hubbard models with spin-orbit coupling are relatively few, particularly as compared to the thoroughly investigated integer fillings, such as $n = 5$ and $n = 4$. To develop a conceptual understanding, here we used doping $n = 3.5$, i.e. 3.5 electrons per site in average, using a model with degenerate bands. Via DMRG calculations here we report the phase diagram varying λ and U . To our best knowledge, theoretical studies at this electronic doping have not been presented before.

The organization of this paper is as follows. In Sec. II, the model used for our study is defined and the details of the computational method are explained. The main results are presented in Sec. III, including the phase diagram varying U and λ . In particular, firstly we present the evidence for the novel block excitonic phase that we unveiled, and then we address the different magnetic phases present in the complete phase diagram, followed by a description of the density of states (DOS). In Sec. IV, we discuss our main results and present our conclusions.

II. MODEL AND METHOD

In this manuscript, we use the three-orbital Hubbard model in the presence of spin-orbit coupling. The Hamiltonian contains a tight-binding term, an on-site Hubbard-Hund interaction, and a spin-orbit coupling: $H = H_K + H_{\text{int}} + H_{\text{SOC}}$. The tight-binding portion is

$$H_K = - \sum_{i,\sigma,\gamma,\gamma'} t_{\gamma\gamma'} (c_{i\sigma\gamma}^\dagger c_{i+1\sigma\gamma'} + \text{h.c.}) + \sum_{i,\sigma,\gamma} \Delta_\gamma n_{i\sigma\gamma}. \quad (1)$$

To gain conceptual understanding, we have focused on the simplest case of degenerate t_{2g} states, hence we fixed $t_{\gamma\gamma'} = t\delta_{\gamma\gamma'}$, where $t = 0.5$, and $\Delta_\gamma = 0$. This leads to a total bandwidth (W) = 2.0 eV in the non-interacting limit. The on-site Hubbard-Hund interaction is

$$H_{\text{int}} = U \sum_{i,\gamma} n_{i\uparrow\gamma} n_{i\downarrow\gamma} + (U' - J_H/2) \sum_{i,\gamma < \gamma'} n_{i\gamma} n_{i\gamma'} - 2J_H \sum_{i,\gamma < \gamma'} \mathbf{S}_{i\gamma} \cdot \mathbf{S}_{i\gamma'} + J_H \sum_{i,\gamma < \gamma'} \left(P_{i\gamma}^\dagger P_{i\gamma'} + \text{h.c.} \right). \quad (2)$$

In the above expression $n_{i\gamma}$ is the electronic density at orbital γ and lattice site i , while the operator $\mathbf{S}_{i\gamma} = \frac{1}{2} \sum_{\alpha,\beta} c_{i\alpha\gamma}^\dagger \sigma_{\alpha\beta} c_{i\beta\gamma}$ is the total spin. The first two terms describe the intra- and inter-orbital electronic repulsion, respectively. The third term contains the Hund coupling that favors the ferromagnetic alignment of the spins at different orbitals; the fourth term is the pair hopping with $P_{i\gamma} = c_{i\downarrow\gamma} c_{i\uparrow\gamma}$ as the pair operator. We assume the standard relation $U' = U - 2J_H$ based on rotational invariance, and we fixed $J_H = U/4$ as in [28]. Hence, only U and λ are free parameters in our study.

The spin-orbit coupling (SOC) term is

$$H_{\text{SOC}} = \lambda \sum_{i,\gamma,\gamma',\sigma,\sigma'} \langle \gamma | \mathbf{L}_i | \gamma' \rangle \cdot \langle \sigma | \mathbf{S}_i | \sigma' \rangle c_{i\sigma\gamma}^\dagger c_{i\sigma'\gamma'}, \quad (3)$$

where λ is the SOC strength.

In the non-interacting limit, both the SOC and tight-binding terms can be diagonalized simultaneously to obtain the following Hamiltonian:

$$H_K + H_{\text{SOC}} = \sum_{k,m} (2t \cos(k) - \frac{\lambda}{2}) a_{k,\frac{3}{2},m}^\dagger a_{k,\frac{3}{2},m} + \sum_{k,m} (2t \cos(k) + \lambda) a_{k,\frac{1}{2},m}^\dagger a_{k,\frac{1}{2},m}. \quad (4)$$

Above we used $a_{k,j_{\text{eff}},m}^\dagger = 1/\sqrt{L} \sum_l e^{-ilk} a_{l,j_{\text{eff}},m}^\dagger$, where $a_{l,j_{\text{eff}},m}^\dagger$ is the creation operator for an electron with total effective angular momentum j_{eff} and z -projection m . The transformation between the t_{2g} orbitals and j_{eff} basis is the following (real-space site index l is dropped):

$$\begin{bmatrix} a_{\frac{3}{2},\frac{3s}{2}} \\ a_{\frac{3}{2},-\frac{s}{2}} \\ a_{\frac{1}{2},-\frac{s}{2}} \end{bmatrix} = \begin{bmatrix} \frac{is}{\sqrt{2}} & \frac{1}{\sqrt{2}} & 0 \\ \frac{s}{\sqrt{6}} & \frac{i}{\sqrt{6}} & \frac{2}{\sqrt{6}} \\ \frac{-s}{\sqrt{3}} & \frac{-i}{\sqrt{3}} & \frac{1}{\sqrt{3}} \end{bmatrix} \begin{bmatrix} c_{\sigma yz} \\ c_{\sigma xz} \\ c_{\bar{\sigma} xy} \end{bmatrix}, \quad (5)$$

where s is $1(-1)$ when σ is $\uparrow(\downarrow)$ and $\bar{\sigma} = -\sigma$ [note that from now onwards to avoid complicated notations, when j_{eff} should be used as subindex, sometimes this quantum number will be simply denoted by j]. Equation (4) is useful to understand the non-interacting limit of the model. As λ is increased, the $j_{\text{eff}} = 3/2$ and $j_{\text{eff}} = 1/2$ bands split. For the doping $n = 3.5$ addressed in this study, in the large λ limit all electrons will be located in the

$j_{\text{eff}} = 3/2$ band, making the $j_{\text{eff}} = 3/2$ band fractionally filled and the $j_{\text{eff}} = 1/2$ band empty. This region is called $j_{\text{eff}} = 3/2$ metal, as discussed in Sec. III C. In [69], we also present the local Coulomb interaction term expressed in the j_{eff} basis. Note that the global $U(1)$ symmetries, corresponding to the conservation of number of particles in the (j, m) states separately, are preserved only for $J_H = 0$ and $U = U'$. Thus, for the case of a finite Hund coupling, as investigated in the present paper, these $U(1)$ symmetries are explicitly broken leading to the possibility of long-range order in some channels.

Because our primary interest is to understand the subtle phases emerging from the competition of the Coulomb interaction, spin-orbit coupling, and kinetic energy, we used the DMRG technique which is numerically exact in one dimension. DMRG can treat the above three terms in the Hamiltonian on equal footing. We solved the above described model for various system lengths, such as $L = 8, 16, 24, 32$, and 48 , fixing the average local density to $n = 3.5$. To reduce the cost of the simulations, we have targetted subspaces of the total $J_z^{\text{eff}} = \sum_i (J_z^{\text{eff}})_i$, which is possible because $[H, J_z^{\text{eff}}] = 0$ for the chosen tight-binding parameters (for details see [28]). For the DMRG process, we used up to 1000 states and the corrected single-site DMRG algorithm [61] with correction $a = 0.001 - 0.008$. We performed 35 to 40 sweeps to gain proper convergence to the ground state properties. After convergence, we calculated the spin structure factor $S(q)$, local occupations $\langle n_{jm} \rangle$, local moments \mathbf{S}_i^2 , \mathbf{L}_i^2 , and $(\mathbf{J}_i^{\text{eff}})^2$, and also the exciton pair-pair correlation $\langle \Delta_{j\tilde{m}}^{\dagger\tilde{m}'}(i) \Delta_{j\tilde{m}'}^{\tilde{m}'}(i') \rangle$ in order to construct the phase diagram. Moreover, we also used the DMRG correction vector method [62] with $L = 16$, as well as the Lanczos algorithm [63] with $L = 4$, to calculate the single-particle DOS.

III. RESULTS

In Fig. 1, we show the phase diagram that we obtained varying U and λ in units of the non-interacting bandwidth W , at a fixed average local electronic density $n = 3.5$. The main result is the presence of a “block excitonic condensate”, accompanied with block magnetic ordering (in a $\uparrow\uparrow\downarrow\downarrow$ pattern). We will discuss this novel phase, and other phases, in the following subsections. Also note that our study is in one dimension. For this reason when we express that in a range of U and λ we are located at a particular phase with particular characteristics, this always has to be interpreted in the sense of dominant power-law decaying correlations, as opposed to true long-range order. However, in the supplementary material we show that the $U(1)$ phase characteristic of other excitonic condensates is explicitly broken in our calculation allowing for long-range order in the block exciton condensate regime to develop, compatible with the numerical results to be shown below.

Before discussing the results in detail, note that we expect the boundaries in the phase diagram Fig. 1 to

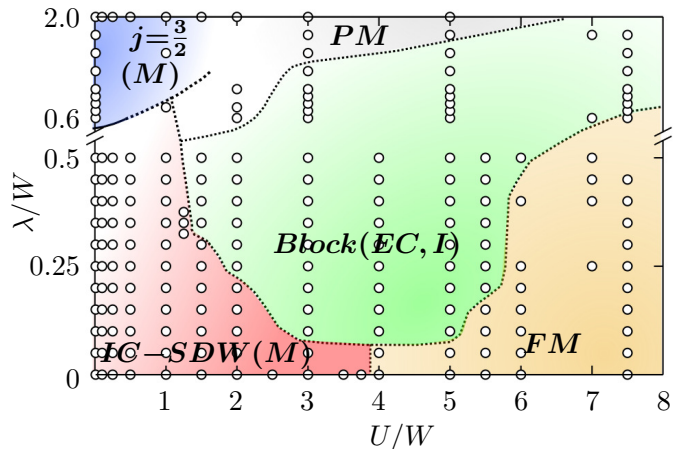


FIG. 1. Shown is the main result of our publication, namely the $\lambda - U$ phase diagram for $n = 3.5$, where $W = 2.0$ eV. The labels $IC - SDW$, $Block(EC)$, FM , and PM stand for Incommensurate Spin Density Wave, Block Excitonic Condensate, Ferromagnetic, and Paramagnetic phases, respectively. $j = 3/2$ is a metallic phase at large SOC where all electrons are in a $j = 3/2$ band, split from the $j = 1/2$ band that is empty. The labels M and I are used in the $IC - SDW$, $Block EC$, and $j = 3/2$ phases to denote metal and insulator, respectively. For the other regimes we could not establish with certainty their metallic vs insulating character. The high density of points was achieved by using a system size $L = 16$ for the DMRG calculations, but several points in the phase diagram were obtained with $L = 32$ chains, as described for special cases below.

correspond to first-order transitions if different symmetries are broken on both sides. In other words, at zero temperature we expect that each boundary between different symmetry breaking patterns corresponds to a level crossing. Computationally this could be tested by plotting the ground state energy varying for example U/W at constant SOC, using a grid of couplings very close to the transition. If a discontinuity in the ground state energy first derivative is found at the transition, a first-order transition occurs. However, these calculations are very time consuming in our multi-orbital model, thus a confirmation of our expectation of first-order transitions is left for future work. Of course, if two regimes are merely separated by a crossover, then this crossover region resembles a continuous transition qualitatively.

A. Condensation of spin-orbit excitons

We now proceed to show and discuss the evidence of excitonic condensation in our phase diagram. We define the creation operator for an exciton at site i as $\Delta_{j\tilde{m}}^{\dagger\tilde{m}'}(i) = a_{ijm}^{\dagger} a_{i\tilde{j}\tilde{m}'}$, where $j = 1/2$ and $\tilde{j} = 3/2$ are fixed. The exciton created by the above operator consists of a hole located at a $\tilde{j} = 3/2$ state with projection m' and an electron with $j = 1/2$ with projection m .

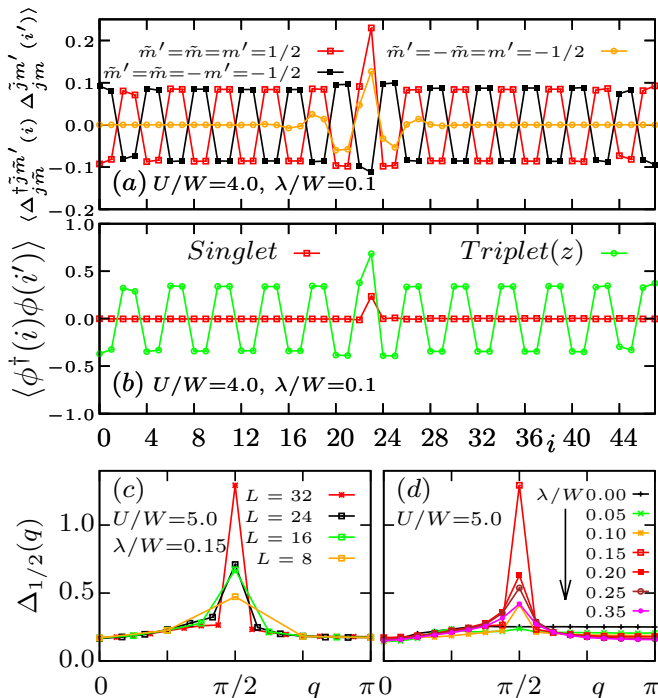


FIG. 2. Panels (a,b) show the exciton-exciton correlation in real space, at $U/W = 4.0$ and $\lambda/W = 0.1$ using a $L = 48$ system size. For panels (a,b), $m = 1/2$ and $i'=23$ (in the middle of a chain with open boundary conditions) are fixed. Panel (c) shows the momentum distribution function for excitons at $U/W = 5.0$ and $\lambda/W = 0.15$. Similar momentum distribution functions of excitons for various λ 's are shown in panel (d). A system size $L = 32$ is used for panel (d).

These excitons are called “spin-orbit excitons” because the electron-hole pair is present in a spin-orbit entangled state. A similar excitonic operator was used before in [25, 28] to investigate the condensation of spin-orbit excitons for the $n = 4$ case. We would like to mention that the present work is the first study where the condensation of these excitons is shown to be stable for $n = 3.5$.

To investigate excitonic condensation we measure the real-space correlation between the excitons i.e. $\langle \Delta_{j\tilde{m}}^{\dagger j\tilde{m}'}(i) \Delta_{j\tilde{m}}^{j\tilde{m}'}(i') \rangle$. Here we would like to mention that earlier similar type of analysis was performed for one-dimensional systems to investigate quasi-excitonic condensations but in simpler models such as the extended Falicov-Kimball model [64].

For the spin-orbit excitons, as the quantum number m can take two values ($m = \pm 1/2$), this gives rise to two channels for excitonic condensation, namely the singlet and triplet channels [29, 65]. We define the exciton cre-

ation operators in both channels in the following manner:

$$\phi^s(i) = \sum_m \Delta_{j\tilde{m}}^{\dagger j\tilde{m}'}(i), \quad (6)$$

$$\phi^t(i) = \sum_{mm'} \Delta_{j\tilde{m}}^{\dagger j\tilde{m}'}(i) \tau_{mm'}, \quad (7)$$

where τ are the Pauli matrices.

In Fig. 2(a), we show the real-space correlations between the exciton pairs (with respect to the central site defined as $i' = 23$) for a system size $L = 48$, at $U/W = 4.0$ and $\lambda/W = 0.1$. In Fig. 2(a), we fix $m = 1/2$. Note the robust block ordering $[+ + - -]$ that we found in the pair-pair correlation between excitons (see “ $\tilde{m}' = \tilde{m} = m' = 1/2$ ”, red line), consisting of holes and electrons with the same projection $j_z^{\text{eff}} = 1/2$. We also noticed the presence of very long distance correlations between excitons of opposite projections (see “ $\tilde{m}' = \tilde{m} = -m' = -1/2$ ” in Fig. 2(a), black line), i.e. one exciton consists of a hole and electron pair with projection $j_z^{\text{eff}} = 1/2$ and the other exciton is made up of a hole and electron with projection $j_z^{\text{eff}} = -1/2$. Note that the above discussed excitonic correlations will contribute to both the singlet and the z component of the triplet channels. Similarly, we can create excitons consisting of electron ($j_z^{\text{eff}} = 1/2$) and hole ($j_z^{\text{eff}} = -1/2$) with different projections: the correlations between these excitons are shown in Fig. 2(a) (see “ $\tilde{m}' = -\tilde{m} = m' = -1/2$ ”, orange line) and they display a rapid exponential decay. These excitonic correlations contribute to the x and y components of the triplet channel.

Using the above information and Eqs.(6,7), we calculate the real-space correlations for the excitons in the singlet and triplet channels (z component). As shown in Fig. 2(b), clearly the triplet channel is the dominant showing quasi long-range order (likely a very slow power-law decay in our finite one dimensional system). As discussed before, the x and y components of the triplet channel have exponential decay. This asymmetry between the x, y , and z components is just a consequence of targeting the total J_z^{eff} sector in our DMRG simulations. For the one-dimensional case studied here, long range correlations (slow power-law decays) is used as evidence for excitonic condensation. For a simpler two-band models, early work [29] showed that condensation of excitons in the triplet channel leads to a spin-density wave and in the singlet channel leads, instead, to a charge-density wave.

Surprisingly, we also observed block magnetic ordering in the excitonic condensate phase reported here, which will be discussed in detail in Sec. III B. The dominating correlations in the z direction of the triplet channel implies that the relevant excitons are created by pairing electron and holes with the same j_z^{eff} , and from now on we will focus only on these excitons.

The momentum distribution function for excitons is $\Delta_m(q) = \frac{1}{L} \sum_{i,i'} \langle \Delta_{j\tilde{m}}^{\dagger j\tilde{m}'}(i) \Delta_{j\tilde{m}}^{j\tilde{m}'}(i') \rangle e^{iq(i-i')}$. This quantity provides an indication of the number of excitons (with projection m) at momentum q . In Fig. 2(d) we

show $\Delta_m(q)$ for $U/W = 5$ and various λ 's. For spin-orbit coupling strength $\lambda/W \lesssim 0.07$, in the ferromagnetic region (to be discussed in more detail in the next subsection), the momentum distribution function is nearly flat. But at larger spin-orbit coupling, excitons condense at momentum $\pi/2$ and on further increasing λ , the number of excitons at $q = \pi/2$ again decreases, as shown in Fig. 2(d). The spin-orbit coupling strength for this crossover from the ferromagnetic phase to the block-excitonic condensate depends on the strength of U . We noticed the interesting feature (see phase diagram in Fig. 1) that for $U/W \gtrsim 5$, a larger U needs a larger λ for the condensation to occur. We also found $\pi/2$ order in the excitonic correlations above the IC-SDW region, as shown in the phase diagram of Fig. 1, contrary to the strong coupling region, where increasing U needs smaller a λ for stabilizing the block-excitonic condensate.

We also show the finite-size scaling of the excitonic momentum distribution function in Fig. 2(c), for system sizes $L = 8, 16, 24$, and 32 . The nearly linear growth of $\Delta_{1/2}(q = \pi/2)$ with the system size (L) again suggests the presence of a robust excitonic condensation. Long-range order is possible because the $U(1)$ symmetry is explicitly broken in the model, as discussed in the supplementary.

B. Magnetic ordering

In this subsection, we will discuss and show the evidence for the different types of magnetic orderings found in the phase diagram. To investigate the various magnetic orderings, we calculate the spin-spin correlation $\langle \mathbf{S}_i \cdot \mathbf{S}_j \rangle$, and associated spin structure factor $S(q) = \frac{1}{L} \sum_{i,j} \langle \mathbf{S}_i \cdot \mathbf{S}_j \rangle e^{i(i-j)q}$. We also calculated the averaged local moments, $\langle \mathbf{S}^2 \rangle = \frac{1}{L} \sum_i \langle \mathbf{S}_i^2 \rangle$. Similarly we calculated $\langle \mathbf{L}^2 \rangle$, $\langle (\mathbf{J}^{\text{eff}})^2 \rangle$, and $\langle \mathbf{L} \cdot \mathbf{S} \rangle$. To evaluate the angle between the average local spin and average local orbital moment, i.e. ϕ_{LS} , we used $\phi_{LS} = \cos^{-1}(\frac{\langle \mathbf{L} \cdot \mathbf{S} \rangle}{\langle l \rangle \langle s \rangle})$, where $\langle l \rangle (\langle l \rangle + 1) = \langle \mathbf{L}^2 \rangle$, and $\langle s \rangle (\langle s \rangle + 1) = \langle \mathbf{S}^2 \rangle$.

For $U/W \gtrsim 3.5$, we found that the IC-SDW region is smoothly connected to the non-interacting limit. In this region (red region in Fig. 1) the local moments gradually form up to the saturated values $\langle \mathbf{S}_i^2 \rangle = 2.875$ and $\langle \mathbf{L}_i^2 \rangle = 1$ as we increase U/W . Eventually this IC-SDW phase crossovers to the ferromagnetic (FM) phase. We noticed that in this IC-SDW region, the spin-ordering vector continuously changes depending on the values of U and λ , as shown in [69], which we believe is the result of Fermi surface renormalization by the combined effect of U and λ . In Fig. 3(a), we show the real-space spin-spin correlation for $U/W = 3$ and $\lambda/W = 0$ depicting the incommensurate order, for a system size $L = 32$. By increasing λ , we found the block magnetic ordering phase above this IC-SDW region, but only as long as $U/W \gtrsim 1.0$, as shown in the phase diagram Fig. 1. This block magnetic order survives up to $\lambda/W \approx 1$. In

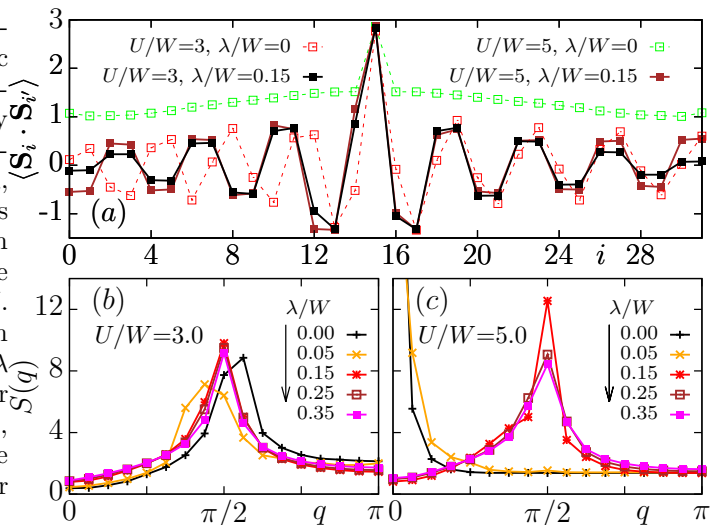


FIG. 3. In panel (a) the real-space spin-spin correlations are shown for $U/W = 3$ and $U/W = 5$, and for $\lambda/W = 0$ and $\lambda/W = 0.15$. In panels (b) and (c), the spin structure factor $S(q)$ is shown for $U/W = 3$ and $U/W = 5$, respectively, and for various values of λ/W , as indicated.

Fig. 3(b), we display the spin structure factor $S(q)$ for $U/W = 3$ for various λ/W 's depicting the clear transition from IC-SDW to block magnetic order. We also show the real-space spin-spin correlation at $U/W = 3$ and $\lambda/W = 0.15$ in Fig. 3(a), portraying the block magnetic order ($\uparrow\uparrow\downarrow\downarrow\uparrow\uparrow$).

As shown in Fig. 1, for $\lambda/W = 0$ the above mentioned IC-SDW phase is directly connected to the ferromagnetic (FM) region in the strong coupling limit; this crossover happens approximately at $U/W \approx 3.9$. In this FM phase, the local spin and orbital moments are fully saturated. The saturated values of the moments can be understood by considering the two-sites case in the large U limit. As we are interested in density $n = 3.5$, in the large U limit the main contribution to the two-sites ground state will arise from the d^3-d^4 configuration. The d^3 site will have local $\langle \mathbf{S}^2 \rangle = \frac{3}{2}(\frac{3}{2} + 1) = 3.75$, while the d^4 site will have local $\langle \mathbf{S}^2 \rangle = 2$, leading to an average 2.875. Similarly the average local moment $\langle \mathbf{L}^2 \rangle = 1$ arises from $\langle \mathbf{L}^2 \rangle = 0$ (for d^3) and $\langle \mathbf{L}^2 \rangle = 2$ (for d^4).

As we increase λ in this FM phase, the system eventually transitions to the block magnetic ordering. In Fig. 3(a), we show the real-space spin-spin correlations for $U/W = 5.0$ and $\lambda/W = 0.0$, and for $U/W = 5.0$ and $\lambda/W = 0.15$, as evidence of the ferromagnetic and block magnetic orders, respectively. The spin structure factors for $U/W = 5.0$ are shown in Fig. 3(c) for various λ 's, depicting the crossover from the ferromagnetic to the block magnetic ordered phases. As discussed in Sec. III A, we suspect this block magnetic ordered phase is related with the condensation of spin-orbit excitons at momentum $q = \pi/2$. Note that for all the points shown in Fig. 1 in the block exciton condensate region

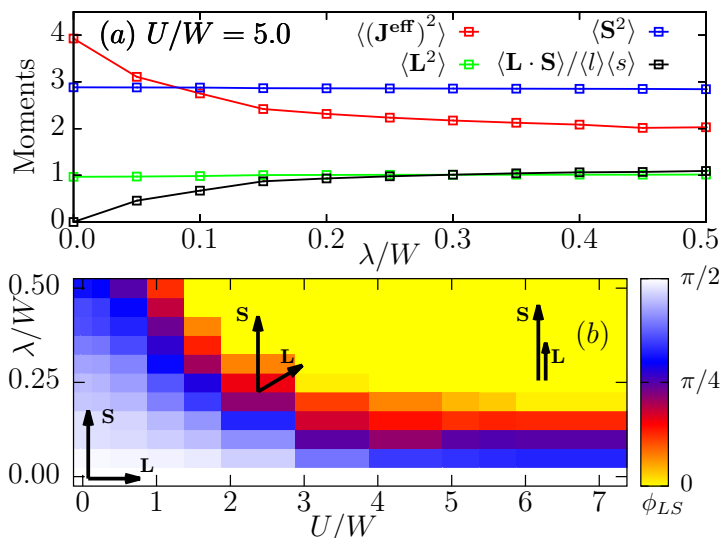


FIG. 4. (a) Average local moments and $\langle \mathbf{L} \cdot \mathbf{S} \rangle / \langle l \rangle \langle s \rangle$ for $U/W = 5$, shown for various λ/W s. In panel (b), ϕ_{LS} is presented for several λ/W 's and U/W 's, where the color depicts the value of ϕ_{LS} as shown in the side panel scale.

(green region in the phase diagram of Fig. 1), we found block magnetic ordering and condensation of excitons at $q = \pi/2$. We also noticed that, as we increase λ , gradually $\Delta_{1/2}(q = \pi/2)$ decreases and the system transitions smoothly into a paramagnetic (PM) phase, with a nearly flat spin structure factor. This PM phase is exotic because it is stabilized at large SOC and large U/W . Its properties will be studied in future work.

Now we turn our focus towards the effect of spin-orbit coupling on the local moments. In Fig. 4(a), we fixed $U/W = 5$, and on increasing λ we observed that the local $\langle \mathbf{S}^2 \rangle$ and $\langle \mathbf{L}^2 \rangle$ remain nearly 2.875 and 1.0, respectively. But although their magnitudes are nearly constant there is a substantial change in the relative orientation of the spin and orbital moments i.e. they gradually modify their relative angle from $\pi/2$ to 0. This rotation affects the local $\langle (\mathbf{J}^{\text{eff}})^2 \rangle$, which decreases as the spin and orbital moments become parallel. In Fig. 4, we show the average angle between the local spin and orbital moments. We notice that, as we increase U , smaller λ 's are enough to render S and L parallel to each other. This indicates that *Coulomb interactions enhance the effect of spin-orbit coupling* and helps to entangle the spin and orbital moments. It is interesting to observe that the novel block excitonic phase we found lies in the region where L and S are parallel to each other with $\langle \mathbf{L} \cdot \mathbf{S} \rangle \approx \langle l \rangle \langle s \rangle \approx 0.8$. We would like to mention that $\langle \mathbf{L} \cdot \mathbf{S} \rangle$ is directly related to the branching ratio calculated by XAS (X-ray Absorption Spectroscopy) experiments for the materials where SOC is robust [66].

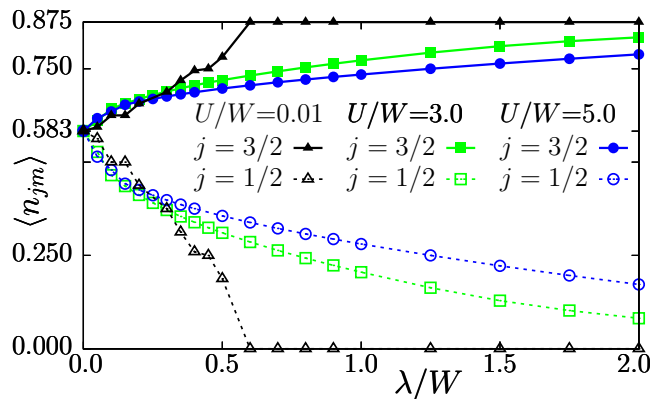


FIG. 5. Average local occupations $\langle n_{jm} \rangle$ shown for $U/W = 0.01$, $U/W = 3.0$, and $U/W = 5.0$, with increasing λ/W . For $j = 1/2$ and $j = 3/2$, m can be $\{\pm 1/2\}$ and $\{\pm 1/2, \pm 3/2\}$, respectively.

C. Local Densities and Density of States

In the last two subsections we established the presence of a novel block excitonic condensate, accompanied with block magnetic order. Now we will discuss the spin-orbit basis-resolved average local occupations $\langle n_{jm} \rangle = \frac{1}{L} \sum_i \langle n_{i,jm} \rangle$ using a $L = 16$ sites system. We also calculate the DOS ($\rho_{jm}(\omega - \mu)$) on a 4-site chain using Lanczos [63], and on a $L = 16$ system using the DMRG correction vector method [62], where μ is the chemical potential evaluated via $(E(N+1) - E(N-1))/2$ for a system with N electrons. In particular, we used the DMRG++ computer program [67] and the Krylov formulation [68] for the DMRG correction vector method [62]. Details on these calculations are provided in [69].

In Fig. 5, we show the effect of spin-orbit coupling on the average local occupations $\langle n_{jm} \rangle$ for three different U values i.e. $U/W = 0.01$, $U/W = 3.0$, and $U/W = 5.0$. Before explaining the results we would like to mention that the occupations in the t_{2g} orbital basis for any λ and U are found to be same i.e. $\langle n_{\sigma\alpha} \rangle \approx 0.5833$, which is consequence of using degenerate orbitals in the kinetic energy term, and Coulomb interaction and spin-orbital coupling that do not break this symmetry in the t_{2g} orbitals. The “good” basis for systems in the presence of spin-orbit coupling (in non-interacting limit) is provided by the j, m states, thus it is reasonable to discuss the occupation in terms of $\langle n_{jm} \rangle$. For small values of Coulomb interaction, such as $U/W = 0.01$, we clearly reproduce the physics of the non-interacting limit. For $\lambda = 0$ we found $\langle n_{jm} \rangle \approx 0.5833$ and as we increase λ the system transitions to a $j_{\text{eff}} = 3/2$ metallic regime where the low-energy $j_{\text{eff}} = 3/2$ band is fractionally filled with $\langle n_{3/2m} \rangle = 0.875$ electrons per site, and the higher energy band $j_{\text{eff}} = 1/2$ is empty. At larger U/W values, this $j_{\text{eff}} = 3/2$ metallic phase is pushed towards larger λ . As also shown in Fig. 5, that for $U/W = 3$ and $U/W = 5$ we do not find a $j_{\text{eff}} = 3/2$ metal for λ/W as large as

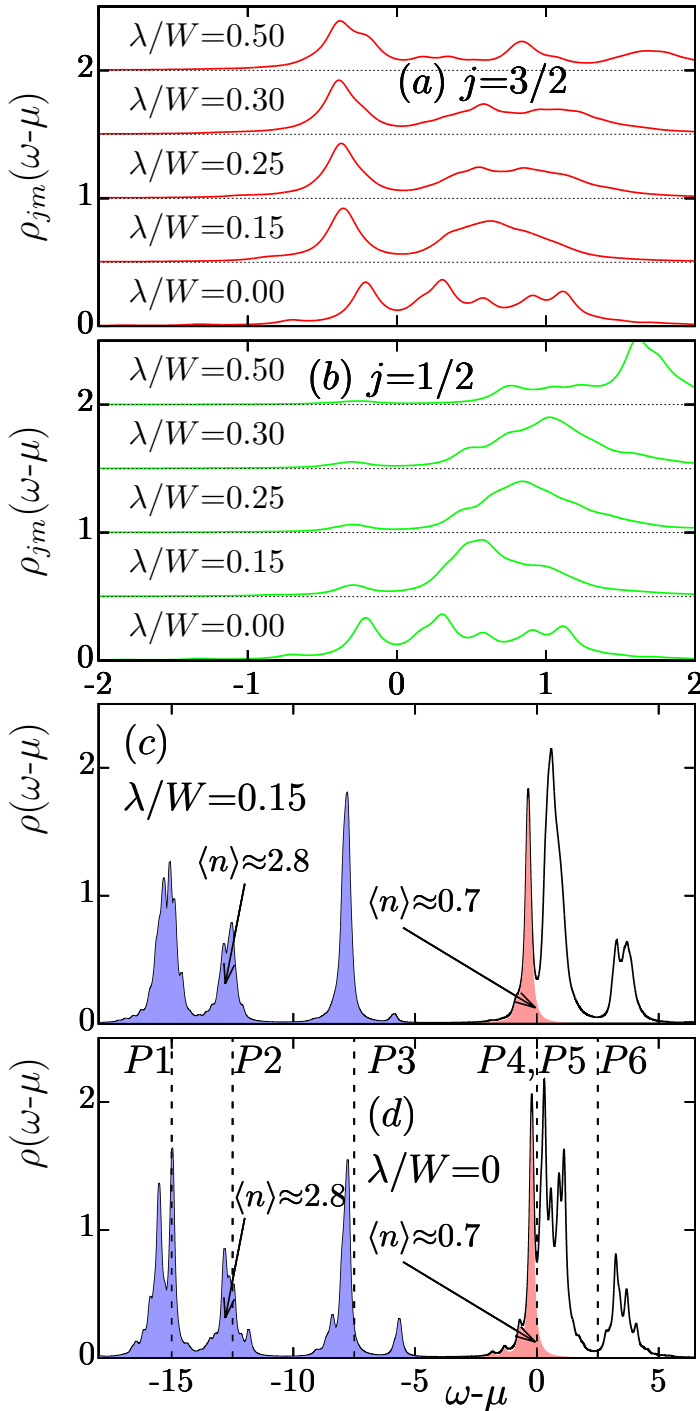


FIG. 6. Density of states, $\rho_{jm}(\omega - \mu)$, near μ shown for various values of λ/W , and at $j_{\text{eff}} = 3/2$ and $j_{\text{eff}} = 1/2$ in panels (a) and (b), respectively. In panels (c) and (d), the total $\rho(\omega - \mu)$ is shown for $\lambda/W = 0.15$ and $\lambda/W = 0$, respectively. All the above results are calculated for a $L = 4$ (OBC) site cluster using Lanczos at $U/W = 5.0$. A broadening $\eta = 0.1$ was used for all the results above. For $j = 1/2$ and $j = 3/2$, m can be $\{\pm 1/2\}$ and $\{\pm 1/2, \pm 3/2\}$, respectively.

2.0. This explains the curvature of the lower boundary

of the $j_{\text{eff}} = 3/2$ metal in the phase diagram Fig. 1. To confirm these results, we also calculated the local occupations for different U/W 's at a fixed $\lambda/W = 2.0$, and found that increasing U gradually increases the filling in the $j_{\text{eff}} = 1/2$ state.

Now let us discuss the DOS calculated using the Lanczos method employing a four-site cluster with open boundary conditions. We checked that even using such a small four-site system, we obtain the same phases as in the phase diagram Fig. 1. Firstly, let us discuss the total DOS $\rho(\omega - \mu) = \sum_{jm} \rho_{jm}(\omega - \mu)$ for $\lambda = 0$, as shown in Fig. 6(d) for $U/W = 5.0$ where we have a FM ground state. We noticed that away from the chemical potential we have four dominant peaks, named $P1$, $P2$, $P3$, and $P6$. These single-particle excitations can be understood in the strong coupling limit using a two-site cluster, as explained in [69]. The interesting feature is the presence of a metallic band near the chemical potential: in the two-site limit this band consists only of two single-particle excitations $P4$ and $P5$. This metallic band in the strong coupling limit [69] contains nearly 0.5 itinerant electrons per site, which are moving in the ferromagnetic background of the other electrons, and leads to a FM metal. For $U/W = 5$, the occupied part of this metallic band is made of nearly 0.7 electrons per site.

Let us investigate the effect of λ on the density of states. In Fig. 6(c), we show $\rho(\omega - \mu)$ for $U/W = 5$ and $\lambda/W = 0.15$. At these couplings our Lanczos results show block magnetic order (with block excitonic order). We noticed that the positions of the peaks away from μ are not changed much, but the DOS at μ decreases with a tendency to open a gap as we move into the block excitonic phase. To explore this issue further, we calculated the j, m -resolved DOS $\rho_{jm}(\omega - \mu)$ near the chemical potential for different λ/W values, as shown in Figs. 6(a,b). For both $j_{\text{eff}} = 3/2$ and $1/2$, note that with increasing λ/W the DOS near μ decreases with the split in the metallic band. But for $j_{\text{eff}} = 1/2$, the DOS below μ gradually decreases to 0, because in the limit of very large λ/W the $j_{\text{eff}} = 1/2$ states will be empty. It should be also noted that for $\lambda/W = 0.5$, we already see the emergence of states near the chemical potential for $j_{\text{eff}} = 3/2$, with the system moving towards the paramagnetic phase. From the above exact analysis of the four-site system, we can conclude that there is a clear tendency towards the opening of a gap in the block phase, thus a clear tendency to form an insulator.

To confirm that the above described results persists for larger systems, we also calculated $\rho_{jm}(\omega - \mu)$ near the chemical potential using the DMRG vector correction method for a $L = 16$ site system. We again fixed $U/W = 5$ and focused on $\lambda/W = 0$ and $\lambda/W = 0.15$, which shows ferromagnetic and block magnetic ordering, respectively, as discussed in Sec. III B. We noted that the metallic band is clearly present in the ferromagnetic phase, see Fig. 7(b). This suggests that indeed there is a fraction of electrons that develop a metallic band, having other localized electrons create a ferromagnetic

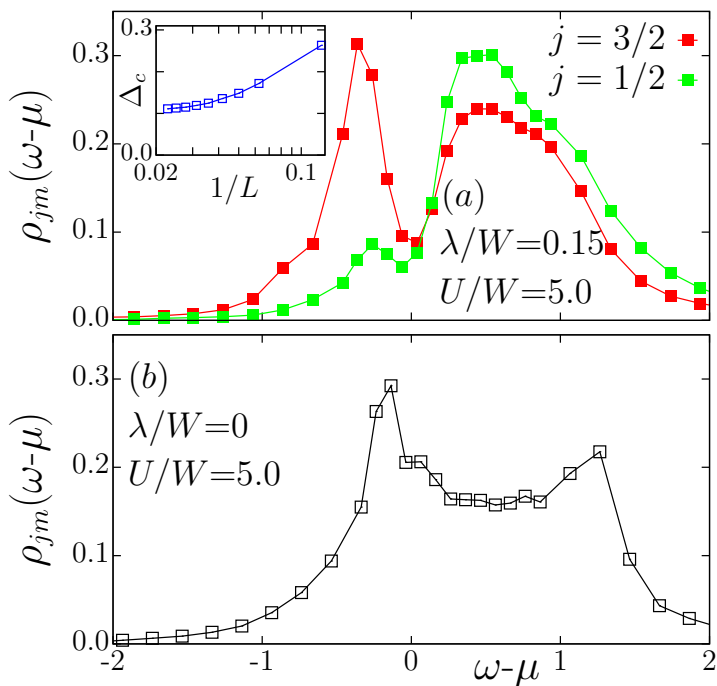


FIG. 7. Density of states, $\rho_{jm}(\omega-\mu)$, shown for a $L = 16$ site system at $\lambda/W = 0.15$ and $\lambda/W = 0$ in panels (a) and (b), respectively. The DMRG vector correction method is used to calculate these results. $U/W = 5.0$ and a broadening $\eta = 0.1$ are employed. The inset in panel (a) contains a finite-size scaling of the charge gap at $\lambda/W = 0.15$ and $U/W = 5.0$. For $j = 1/2$ and $j = 3/2$, m can be $\{\pm 1/2\}$ and $\{\pm 1/2, \pm 3/2\}$, respectively.

background with spins $S \approx 3/2$ [69]. If now we increase λ/W to 0.15, driving the system towards the block excitonic phase, both the $j_{\text{eff}} = 3/2$ and $1/2$ sectors show a tendency to open a gap and being insulating. These results further confirm the understanding deduced from the small $L = 4$ exact result. To establish the insulating nature of the block excitonic phase, we calculated the single-particle charge gap $\Delta_c = E_{N+1} + E_{N-1} - 2E_N$, where N is the total number of electrons. We performed finite-size scaling of the gap Δ_c using various system sizes between $L = 8$ and $L = 44$. As shown in the inset of Fig. 7(a), we noticed that $\lim_{L \rightarrow \infty} \Delta_c(L)$ converges approximately to 0.1 eV.

The novel block phase shown in this publication resembles the block magnetic order phase reported earlier in the context of three-orbital Hubbard models. In fact, block magnetic ordering, without the excitonic condensate component, has been found previously in models without spin-orbit coupling, in the context of the Orbital Selective Mott Phase (OSMP) [70–74]. In the latter, two orbitals are metallic with fractional filling and one orbital is insulating with half-filling. However, note that the block phase discussed in this publication is *not* accompanied by an OSMP phase, as indicated by the av-

erage local occupations. Instead the novel block phase discussed here is accompanied by the condensation of spin-orbit excitons at momentum $q = \pi/2$ for which a finite spin-orbit coupling is a necessary condition.

IV. CONCLUSIONS

In this paper, we used an accurate numerical technique, DMRG, to construct the λ vs U phase diagram for the one-dimensional three-orbital Hubbard model at $n = 3.5$. As our main result, we provide the first numerical evidence for the condensation of spin-orbit excitons in the fractionally filled three-orbital Hubbard model. Our calculations show that the spin-orbit excitons condense in the triplet channel and at momentum $\pi/2$, for all the points shown inside the green region of the phase diagram displayed in Fig. 1. This quasi-condensation of excitons is accompanied by tendencies to open a gap at the chemical potential and also by block magnetic ordering. Interestingly, the block excitonic condensate unveiled here can be stabilized by introducing the spin-orbit coupling on both the IC-SDW and Ferromagnetic metallic phases. We also noticed that in this novel block excitonic condensate phase, local spin and orbital moments are highly entangled and nearly parallel to each other.

We believe the results reported in this publication – which are unique given the considerable computational effort needed that requires vast computational resources – will encourage further theoretical and experimental investigations on fractionally-filled iridates [75–79] and also on other quasi-one dimensional materials with large spin-orbit coupling. While our model calculations cannot establish which precise material will realize the novel phase unveiled, we believe from now on the block condensate has to be considered among the candidate states when $n = 3.5$ materials are studied.

V. ACKNOWLEDGMENTS

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- ¹ G. Cao and P. Schlottmann, [Reports on Progress in Physics, Volume 81, Number 4](#).
- ² R. J. Cava, B. Batlogg, K. Kiyono, H. Takagi, J. J. Krajewski, W. F. Peck, Jr., L. W. Rupp, Jr., and C. H. Chen, [Phys. Rev. B 49, 11890 \(1994\)](#).
- ³ T. Shimura, Y. Inaguma, T. Nakamura, M. Itoh, and Y. Morii, [Phys. Rev. B 52, 9143 \(1995\)](#).
- ⁴ G. Cao, J. Bolivar, S. McCall, J. E. Crow, and R. P. Guertin, [Phys. Rev. B 57, R11039\(R\) \(1998\)](#).
- ⁵ H. Okabe, M. Isobe, E. Takayama-Muromachi, A. Koda, S. Takeshita, M. Hiraiishi, M. Miyazaki, R. Kadono, Y. Miyake, and J. Akimitsu, [Phys. Rev. B 83, 155118 \(2011\)](#).
- ⁶ W. Witczak-Krempa, G. Chen, Y. B. Kim, and L. Balents, [Annu. Rev. Condens. Matter Phys. 5, 57 \(2014\)](#).
- ⁷ H. Watanabe, T. Shirakawa, and S. Yunoki, [Phys. Rev. B 89, 165115 \(2014\)](#).
- ⁸ J. G. Rau, E. K. Lee, and H. Kee, [Annu. Rev. Condens. Matter Phys. 7, 195 \(2016\)](#).
- ⁹ D. Haskel, G. Fabbris, M. Zhernenkov, P. P. Kong, C. Q. Jin, G. Cao, and M. van Veenendaal, [Phys. Rev. Lett. 109, 027204 \(2012\)](#).
- ¹⁰ B. J. Kim, H. Ohsumi, T. Komesu, S. Sakai, T. Morita, H. Takagi, and T. Arima, [Science 323, 1329 \(2009\)](#).
- ¹¹ H. Watanabe, T. Shirakawa, and S. Yunoki, [Phys. Rev. Lett. 105, 216410 \(2010\)](#).
- ¹² B. J. Kim, H. Jin, S. J. Moon, J.-Y. Kim, B.-G. Park, C. S. Leem, J. Yu, T. W. Noh, C. Kim, S.-J. Oh, J.-H. Park, V. Durairaj, G. Cao, and E. Rotenberg, [Phys. Rev. Lett. 101, 076402 \(2008\)](#).
- ¹³ S. Fujiyama, H. Ohsumi, T. Komesu, J. Matsuno, B. J. Kim, M. Takata, T. Arima, and H. Takagi, [Phys. Rev. Lett. 108, 247212 \(2012\)](#).
- ¹⁴ G. Jackeli and G. Khaliullin, [Phys. Rev. Lett. 102, 017205 \(2009\)](#).
- ¹⁵ B. J. Kim, J. Yu, H. Koh, I. Nagai, S. I. Ikeda, S.-J. Oh, and C. Kim, [Phys. Rev. Lett. 97, 106401 \(2006\)](#).
- ¹⁶ C. Martins, M. Aichhorn, L. Vaugier, and S. Biermann, [Phys. Rev. Lett. 107, 266404 \(2011\)](#).
- ¹⁷ B. F. Phelan, E. M. Seibel, D. Badoe, Jr., W. Xie, and R. Cava, [Solid State Commun. 236, 37 \(2016\)](#).
- ¹⁸ D. C. Johnston, [Adv. Phys. 59, 803 \(2010\)](#).
- ¹⁹ P. C. Dai, J. P. Hu, and E. Dagotto, [Nat. Phys. 8, 709 \(2012\)](#), and references therein.
- ²⁰ D. Liu, W. Zhang, D. Mou, J. He, Y. Ou, Q. Wang, Z. Li, L. Wang, L. Zhao, S. He et al., [Nat. Commun. 3, 931 \(2012\)](#).
- ²¹ S. V. Borisenko, D. V. Evtushinsky, Z.-H. Liu, I. Morozov, R. Kappenberger, S. Wurmehl, B. Büchner, A. N. Yaresko, T. K. Kim, M. Hoesch, T. Wolf, and N. D. Zhigadlo, [Nat. Phys. 12, 311 \(2016\)](#).
- ²² G. Khaliullin, [Phys. Rev. Lett. 111, 197201 \(2013\)](#).
- ²³ O. N. Meetei, W. S. Cole, M. Randeria, and N. Trivedi, [Phys. Rev. B 91, 054412 \(2015\)](#).
- ²⁴ C. Svoboda, M. Randeria, and N. Trivedi, [Phys. Rev. B 95, 014409 \(2017\)](#).
- ²⁵ T. Sato, T. Shirakawa, and S. Yunoki, [Phys. Rev. B 99, 075117 \(2019\)](#).
- ²⁶ T. Sato, T. Shirakawa, and S. Yunoki, [Phys. Rev. B 91, 125122 \(2015\)](#).
- ²⁷ A. J. Kim, H. O. Jeschke, P. Werner, and R. Valentí, [Phys. Rev. Lett. 118, 086401 \(2017\)](#).
- ²⁸ N. Kaushal, J. Herbrych, A. Nocera, G. Alvarez, A. Moreo, F. A. Reboredo, E. Dagotto, [Phys. Rev. B 96, 155111 \(2017\)](#).
- ²⁹ D. Jérôme, T. M. Rice, and W. Kohn, [Phys. Rev. 158, 462 \(1967\)](#); B. I. Halperin and T. M. Rice, [Rev. Mod. Phys. 40, 755 \(1968\)](#); B. I. Halperin and T. M. Rice, [Solid State Physics 21, 115 \(1968\)](#).
- ³⁰ G. Cao, T. F. Qi, L. Li, J. Terzic, S. J. Yuan, L. E. DeLong, G. Murthy, and R. K. Kaul, [Phys. Rev. Lett. 112, 056402 \(2014\)](#).
- ³¹ L. T. Corredor, G. Aslan-Cansever, M. Sturza, Kaustuv Manna, A. Maljuk, S. Gass, T. Dey, A. U. B. Wolter, Olga Kataeva, A. Zimmermann, M. Geyer, C. G. F. Blum, S. Wurmehl, and B. Büchner, [Phys. Rev. B 95, 064418 \(2017\)](#).
- ³² J. Terzic, H. Zheng, F. Ye, H. D. Zhao, P. Schlottmann, L. De Long, and G. Cao, [Phys. Rev. B 96, 064436 \(2017\)](#).
- ³³ T. Dey, A. Maljuk, D. V. Efremov, O. Kataeva, S. Gass, C. G. F. Blum, F. Steckel, D. Gruner, T. Ritschel, A. U. B. Wolter, J. Geck, C. Hess, K. Koepernik, J. van den Brink, S. Wurmehl, and B. Buchner, [Phys. Rev. B 93, 014434 \(2016\)](#).
- ³⁴ A. Nag, S. Bhowal, A. Chakraborty, M. M. Sala, A. Efimenko, F. Bert, P. K. Biswas, A. D. Hillier, M. Itoh, S. D. Kaushik, V. Siruguri, C. Meneghini, I. Dasgupta, and S. Ray, [Phys. Rev. B 98, 014431 \(2018\)](#).
- ³⁵ M. Kusch, V. M. Katukuri, N. A. Bogdanov, B. Buchner, T. Dey, D. V. Efremov, J. E. Hamann-Borrero, B. H. Kim, M. Krisch, A. Maljuk, M. Moretti Sala, S. Wurmehl, G. Aslan-Cansever, M. Sturza, L. Hozoi, J. van den Brink, and J. Geck, [Phys. Rev. B 97, 064421 \(2018\)](#).
- ³⁶ M. Souri, B. H. Kim, J. H. Gruenewald, J. G. Connell, J. Thompson, J. Nichols, J. Terzic, B. I. Min, G. Cao, J. W. Brill, and A. Seo, [Phys. Rev. B 95, 235125 \(2017\)](#).
- ³⁷ J. Kim, D. Casa, M. H. Upton, T. Gog, Y.-J. Kim, J. F. Mitchell, M. van Veenendaal, M. Daghofer, J. van den Brink, G. Khaliullin, and B. J. Kim, [Phys. Rev. Lett. 108, 177003 \(2012\)](#).
- ³⁸ J. Kim, M. Daghofer, A. H. Said, T. Gog, J. van den Brink, G. Khaliullin, and B. J. Kim, [Nat. Comm. 5, 4453 \(2014\)](#).
- ³⁹ J. H. Gruenewald, J. Kim, H. S. Kim, J. M. Johnson, J. Hwang, M. Souri, J. Terzic, S. H. Chang, A. Said, J. W. Brill, G. Cao, H. Y. Kee, and S. S. A. Seo, [Adv. Mater. 29, 1603798 \(2016\)](#).
- ⁴⁰ N. A. Bogdanov, V. M. Katukuri, H. Stoll, J. van den Brink, and L. Hozoi, [Phys. Rev. B 85, 235147 \(2012\)](#).
- ⁴¹ K. Ohgushi, J.-I. Yamaura, H. Ohsumi, K. Sugimoto, S. Takeshita, A. Tokuda, H. Takagi, M. Takata, and T.-H. Arima, [Phys. Rev. Lett. 110, 217212 \(2013\)](#).
- ⁴² M. Moretti Sala, K. Ohgushi, A. Al-Zein, Y. Hirata, G. Monaco, and M. Krisch, [Phys. Rev. Lett. 112, 176402 \(2014\)](#).
- ⁴³ S.-W. Kim, C. Liu, H.-J. Kim, J.-H. Lee, Y. Yao, K.-M. Ho, and J.-H. Cho, [Phys. Rev. Lett. 115, 096401 \(2015\)](#).
- ⁴⁴ G. Cao, V. Durairaj, S. Chikara, S. Parkin, and P. Schlottmann, [Phys. Rev. B 75, 134402 \(2007\)](#).
- ⁴⁵ S. Calder, G.-X. Cao, S. Okamoto, J. W. Kim, V. R. Cooper, Z. Gai, B. C. Sales, M. D. Lumsden, D. Mandrus, and A. D. Christianson, [Phys. Rev. B 89, 081104\(R\) \(2014\)](#).

- ⁴⁶ K. Maiti, R. S. Singh, V. R. R. Medicherla, S. Rayaprol, and E. V. Sampathkumaran, *Phys. Rev. Lett.* **95**, 016404 (2005).
- ⁴⁷ O. B. Korneta, S. Chikara, S. Parkin, L. E. DeLong, P. Schlottmann, and G. Cao, *Phys. Rev. B* **81**, 045101 (2010).
- ⁴⁸ M. Ye, H-S. Kim, J-W. Kim, C-J. Won, K. Haule, D. Vanderbilt, S-W. Cheong, and G. Blumberg, *Phys. Rev. B* **98**, 201105(R) (2018).
- ⁴⁹ X. Liu, V. M. Katukuri, L. Hozoi, W.-G. Yin, M. P. M. Dean, M. H. Upton, J. Kim, D. Casa, A. Said, T. Gog, T. F. Qi, G. Cao, A. M. Tsvelik, J. van den Brink, and J. P. Hill, *Phys. Rev. Lett.* **109**, 157401 (2012).
- ⁵⁰ W-G. Yin, X. Liu, A. M. Tsvelik, M. P. M. Dean, M. H. Upton, JungHo Kim, D. Casa, A. Said, T. Gog, T. F. Qi, G. Cao, and J. P. Hill, *Phys. Rev. Lett.* **111**, 057202 (2013).
- ⁵¹ R. Morrow, M. A. Susner, M. D. Sumption, and P. M. Woodward, *Phys. Rev. B* **92**, 134402 (2015).
- ⁵² J. Terzic, J. C. Wang, Feng Ye, W. H. Song, S. J. Yuan, S. Aswartham, L. E. DeLong, S. V. Streltsov, D. I. Khomskii, and G. Cao, *Phys. Rev. B* **91**, 235147 (2015).
- ⁵³ S. V. Streltsov, G. Cao, and D. I. Khomskii, *Phys. Rev. B* **96**, 014434 (2017).
- ⁵⁴ Y. Wang, R. Wang, J. Kim, M. H. Upton, D. Casa, T. Gog, G. Cao, G. Kotliar, M. P. M. Dean, and X. Liu, *ArXiv:1810.05243* (2018).
- ⁵⁵ X. Chen and S. D. Wilson, *Phys. Rev. B* **94**, 195115 (2016).
- ⁵⁶ K. Matsuhira, K. Nakamura, Y. Yasukuni, Y. Yoshimoto, D. Hirai, and Z. Hiroi, *J. Phys. Soc. Jpn.* **87**, 013703 (2018).
- ⁵⁷ Z. Q. Mao, T. He, M. M. Rosario, K. D. Nelson, D. Okuno, B. Ueland, I. G. Deac, P. Schiffer, Y. Liu, and R. J. Cava, *Phys. Rev. Lett.* **90**, 186601 (2003).
- ⁵⁸ W. Kobayashi, *Phys. Rev. B* **79**, 155116 (2009).
- ⁵⁹ S. R. White and R. M. Noack, *Phys. Rev. Lett.* **68**, 3487 (1992).
- ⁶⁰ S. R. White, *Phys. Rev. B* **48**, 10345 (1993).
- ⁶¹ S. R. White, *Phys. Rev. B* **72**, 180403(R) (2005).
- ⁶² T. Kühner and S. R. White, *Phys. Rev. B* **60**, 335 (1999).
- ⁶³ E. Dagotto, *Rev. Mod. Phys.* **66**, 763 (1994), and references therein.
- ⁶⁴ S. Ejima, T. Kaneko, Y. Ohta, and H. Fehske, *Phys. Rev. Lett.* **112**, 026401(2014)
- ⁶⁵ J. Kuneš, *Journal of Physics: Condensed Matter*, Volume **27**, Number 33 (2015).
- ⁶⁶ M. A. Laguna-Marco, D. Haskel, N. Souza-Neto, J. C. Lang, V. V. Krishnamurthy, S. Chikara, G. Cao, and M. van Veenendaal, *Phys. Rev. Lett.* **105**, 216407 (2010).
- ⁶⁷ G. Alvarez, *Computer Physics Communications* **180**, 1572 (2009).
- ⁶⁸ A. Nocera and G. Alvarez, *Phys. Rev. E* **94**, 053308 (2016)
- ⁶⁹ See the supplementary material.
- ⁷⁰ J. Rincón, A. Moreo, G. Alvarez, and E. Dagotto, *Phys. Rev. Lett.* **112**, 106405 (2014).
- ⁷¹ J. Rincón, A. Moreo, G. Alvarez, and E. Dagotto, *Phys. Rev. B* **90**, 241105(R) (2014).
- ⁷² G. Liu, N. Kaushal, S. Li, C. B. Bishop, Y. Wang, S. Johnston, G. Alvarez, A. Moreo, and E. Dagotto, *Phys. Rev. E* **93**, 063313 (2016).
- ⁷³ S. Li, N. Kaushal, Y. Wang, Y. Tang, G. Alvarez, A. Nocera, T. A. Maier, E. Dagotto, and S. Johnston, *Phys. Rev. B* **94**, 235126 (2016).
- ⁷⁴ J. Herbrych, N. Kaushal, A. Nocera, G. Alvarez, A. Moreo, and E. Dagotto, *Nat. Comm.* **9**, 3736 (2018).
- ⁷⁵ J. Gunasekera, Y. Chen, J. W. Kremenak, P. F. Miceli, and D. K. Singh, *Journal of Physics: Condensed Matter*, Volume **27**, Number 5 (2015).
- ⁷⁶ J. Gunasekera, L. Harriger, A. Dahal, T. Heitmann, G. Vignale, and D. K. Singh, *Scientific Reports* volume **5**, Article number: 18047 (2015).
- ⁷⁷ W. Ju, G-Q. Liu, and Z. Yang, *Phys. Rev. B* **87**, 075112 (2013).
- ⁷⁸ J. Chakraborty, *Phys. Rev. B* **97**, 235147 (2018).
- ⁷⁹ S. K. Panda, S. Bhowal, Y. Li, S. Ganguly, R. Valenti, L. Nordstrom, and I. Dasgupta, *Phys. Rev. B* **92**, 180403(R) (2015).