

Coherent Quasiparticles with a Small Fermi Surface in Lightly Doped $\text{Sr}_3\text{Ir}_2\text{O}_7$

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We characterize the electron doping evolution of $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ by means of angle-resolved photoemission. Concomitant with the metal insulator transition around $x \approx 0.05$ we find the emergence of coherent quasiparticle states forming a closed small Fermi surface of volume $3x/2$, where x is the independently measured La concentration. The quasiparticle weight Z remains large along the entire Fermi surface, consistent with the moderate renormalization of the low-energy dispersion, and no pseudogap is observed. This indicates a conventional, weakly correlated Fermi liquid state with a momentum independent residue $Z \approx 0.5$ in lightly doped $\text{Sr}_3\text{Ir}_2\text{O}_7$.

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The 5d transition metal iridium oxides may host exotic quantum phases emerging from the interplay of correlations and strong spin-orbit coupling [1,2]. Iridates of the Ruddlesden-Popper series $\text{Sr}_{n+1}\text{Ir}_n\text{O}_{3n+1}$ share key structural and electronic properties with the parent compounds of copper oxide superconductors [3–5] and are thus of particular interest as candidate materials for engineering unconventional superconductivity [6,7]. Despite their partially filled 5d shell with a single hole per Ir in the t_{2g} manifold, the layered $n = 1, 2$ members are antiferromagnetic insulators. This has been attributed to a reduced orbital degeneracy and one-electron bandwidth resulting from strong spin-orbit coupling and structural distortions leading to a single, narrow $j_{\text{eff}} = 1/2$ band that is susceptible to the moderate electron correlations in the Ir t_{2g} shell [3,8,9]. This picture is supported by band structure calculations for single layer Sr_2IrO_4 . Its low-energy electronic structure can indeed be approximated by a single $j_{\text{eff}} = 1/2$ band, which opens a gap in the presence of electron correlations [10,11]. The resulting insulating state shows in-plane ordered moments and the characteristic spin dynamics of a Heisenberg antiferromagnet with a gapless excitation spectrum [4,5,12,13]. The effective low-energy physics can thus be described in a Hubbard model with exchange interaction comparable to cuprates but an electronlike noninteracting Fermi surface centered at Γ [4,7]. Intriguingly, a recent Monte Carlo study of this Hamiltonian predicts d -wave superconductivity for

electron doping but not for hole doping [7]. Moreover, recent angle-resolved photoemission (ARPES) data from lightly doped Sr_2IrO_4 [14,15] reproduced much of the unique phenomenology observed in underdoped cuprates, including open Fermi arcs and a temperature dependent pseudogap [16–18]. Evidence for strong correlations were also reported for Ru-doped $\text{Sr}_3\text{Ir}_2\text{O}_7$ [19].

While the effective single band pseudospin-1/2 Mott picture is well established for Sr_2IrO_4 [10,11], the properties of the bilayer compound are more ambiguous. On one hand, the magnetic structure of $\text{Sr}_3\text{Ir}_2\text{O}_7$ provides evidence for a $j_{\text{eff}} = 1/2$ state. Both the out-of-plane collinear ordering and the large spin-wave gap follow naturally from the pseudodipolar interaction characteristic of the $j_{\text{eff}} = 1/2$ state [20–22]. Moreover, the ordered moment of $\text{Sr}_3\text{Ir}_2\text{O}_7$ is comparable to Sr_2IrO_4 [23]. On the other hand, band structure calculations show mixing of $j_{\text{eff}} = 1/2$ and $3/2$ states in $\text{Sr}_3\text{Ir}_2\text{O}_7$ and a semimetallic electronic structure with a direct band gap throughout the Brillouin zone but band overlap from different k points, qualitatively different from the case of Sr_2IrO_4 [10]. Experimental evidence for a departure from the ideal $j_{\text{eff}} = 1/2$ Mott state comes from the small, possibly indirect, charge gap [24–26] and the significant overlap in energy between $j_{\text{eff}} = 1/2$ and $3/2$ states found in previous ARPES studies on undoped $\text{Sr}_3\text{Ir}_2\text{O}_7$ [27–30]. A paramagnetic metallic ground state was reported in $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ for La concentrations above $x \approx 0.03$ [31]. Yet, little is known

about the microscopic electronic structure of doped $\text{Sr}_3\text{Ir}_2\text{O}_7$.

Here we show that lightly doped $\text{Sr}_3\text{Ir}_2\text{O}_7$ supports a weakly correlated Fermi-liquid-like metallic state, strikingly different from the electronic structure of underdoped cuprates and from recent ARPES results from doped Sr_2IrO_4 [14,15].

ARPES measurements were performed at the I05 beam line of the Diamond Light Source. Crystals of $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ were flux grown and showed a paramagnetic metallic ground state for $x > 0.045$. Details of the sample growth and characterization will be presented elsewhere [32]. The samples were cleaved at pressures $< 10^{-10}$ mbar and temperatures < 50 K. Measurements were made using photon energies between 30 and 120 eV. All presented data were acquired at 83 eV with an energy resolution of 15 meV. The sample temperature was 8 and 50 K for conducting and insulating samples, respectively. The electronic structure calculations were performed within the local density approximation (LDA) using the WIEN2k software package [33]. The effect of spin-orbit interaction was included using a second-variational scheme. Doping was treated within the virtual crystal approximation (VCA). We used the $Bbcb$ space group with lattice parameters $a = 5.517$, $b = 5.515$, $c = 20.897$ Å [32] and relaxed the internal atomic positions for each value of doping. In our LDA + SO + U calculations we use $U = 2$ eV and the double counting prescription of Ref. [34]. All Ir atoms are kept equivalent so that the antiferromagnetic solution is not allowed by symmetry and the calculations converge to a paramagnetic state.

Figure 1 shows the main features of the low energy electronic structure of electron doped $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ for $x = 0.065$ where the samples are metallic down to low temperatures [31,32]. The spectral weight at the Fermi level is concentrated along the Brillouin zone diagonal and

strongly suppressed along the Ir-Ir nearest neighbor direction, reminiscent of the nodal-antinodal dichotomy in lightly doped cuprates [17,18] and the recent observation of Fermi arcs in doped Sr_2IrO_4 [14,15]. Despite not having any special meaning for the iridates, we use the term “nodal” throughout this Letter to facilitate the comparison with the copper oxides. The dispersion plots and energy distribution curves (EDCs) shown in Figs. 1(b)–1(d) confirm this picture. We find coherent quasiparticlelike states along the nodal cut but only faint low-energy spectral weight with a broader momentum distribution along the antinodal direction ΓX . The intense holelike bands observed at higher energy (-0.3 eV at Γ and -0.05 eV at X) originate from the occupied $j_{\text{eff}} = 3/2$ and $1/2$ states in the undoped parent compound, respectively [11,28]. Along the nodal direction, where the quasiparticle states are clearly separated from the bands of the parent compound, we observe incoherent tails with nearly vertical dispersion separated from the quasiparticle bands by a slight suppression of spectral weight resulting in a peak-dip-hump line shape indicative of coupling to a bosonic mode. While the distribution of spectral weight alone does not permit a reliable determination of the coupling strength, the dominant mode energy can be estimated from the separation of the peak and hump to be around 100–150 meV and thus in the range where the magnon density of states peaks [22].

The systematic evolution of the electronic structure of $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ with increasing doping is shown in Fig. 2. For $x = 0.01$, faint spectral weight appears around the M point in the gap of the parent compound, indicating localized carriers, consistent with the insulating ground state of these samples. At $x = 0.05$, coincident with the metal insulator transition in transport, we observe the emergence of coherent quasiparticle states whose weight, Fermi wave vector, and bandwidth increase monotonically with doping. We note that the doping level at which the first

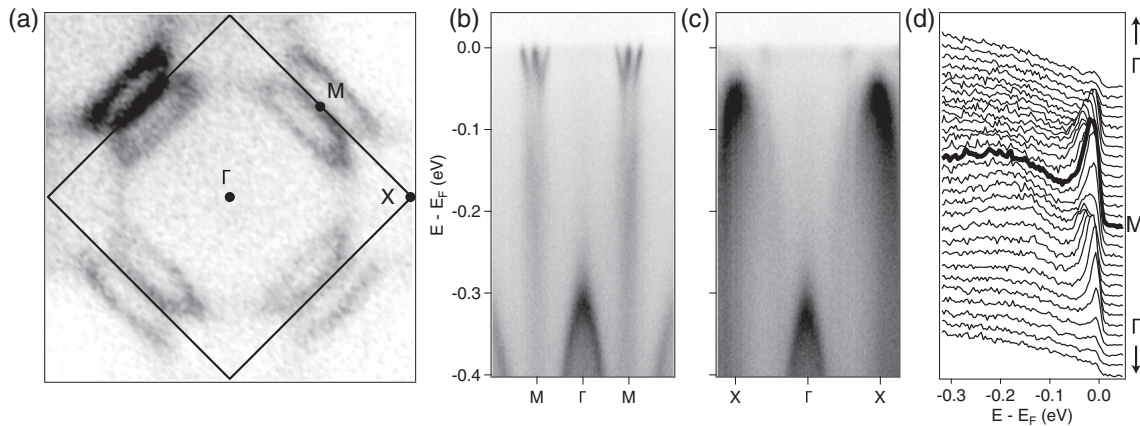


FIG. 1. Low energy excitations in lightly doped $\text{Sr}_3\text{Ir}_2\text{O}_7$. (a) Experimental Fermi surface of $(\text{Sr}_{0.935}\text{La}_{0.065})_3\text{Ir}_2\text{O}_7$ taken with $h\nu = 83$ eV at 8 K. We denote the $(\pi, 0)$ point of the two-dimensional Ir sublattice by X and the $(\pi/2, \pi/2)$ point by M . (b), (c) Photoemission intensity along the ΓM and ΓX high symmetry directions, respectively. (d) Stack of energy distribution curves around the Fermi surface extracted from the data shown in panel (b).

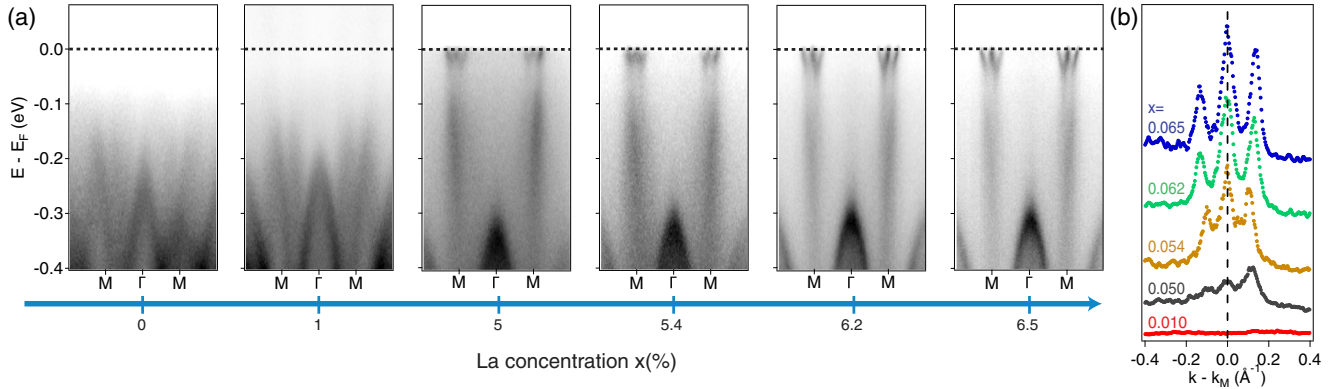


FIG. 2 (color online). Doping evolution of $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$. (a) Dispersion along the nodal ΓM direction. Coherent quasiparticle states emerge for $x > 0.05$. (b) momentum distribution curves (MDCs) at the Fermi energy. The Fermi wave vector increases monotonically with doping, indicative of an increasing density of itinerant carriers.

quasiparticlelike states appear in $\text{Sr}_3\text{Ir}_2\text{O}_7$ and their nodal Fermi velocity of $\approx 10^5$ m/s are comparable to lightly doped cuprates [17,35–37]. However, the quasiparticle pole is more pronounced here and more clearly separated from the incoherent weight at higher energy.

For the remainder of this Letter, we discuss the origin of the phenomenology described above. To this end, we first compare our data with band structure calculations. Using LDA + SO calculations, we find a metallic ground state of undoped $\text{Sr}_3\text{Ir}_2\text{O}_7$, consistent with earlier reports [24,25]. The doubling of the in-plane unit cell due to the rotations of the IrO_6 octahedra around the c axis and the bilayer nature of $\text{Sr}_3\text{Ir}_2\text{O}_7$ causes an even number of Ir $5d^5$ ions per unit cell and thus an even number of electrons. Yet, the significant itinerancy of the states prevents the formation of a band insulator and leads to a compensated metallic state with a small hole pocket at Γ and electron pockets at the M points. However, there is a direct gap between the bands that give rise to the hole and electron pockets (see the Supplemental Material [38]) rendering the LDA + SO band structure of $\text{Sr}_3\text{Ir}_2\text{O}_7$ characteristic of a semimetal, clearly distinct from Sr_2IrO_4 [10]. The hole pocket in $\text{Sr}_3\text{Ir}_2\text{O}_7$ is rapidly pushed below the Fermi level by adding correlations within paramagnetic LDA + SO + U , leading to an insulating ground state [24,25]. Thus, in this description, undoped $\text{Sr}_3\text{Ir}_2\text{O}_7$ is an insulator whose gap is underestimated within LDA + SO.

In Fig. 3 we superimpose LDA + SO + U Fermi surfaces for different doping levels treated within the VCA on data taken for $x = 0.065$. The good agreement of these calculations with the data strongly suggests that the nodal-antinode dichotomy is a band structure effect and thus of fundamentally different origin than in the cuprates. This interpretation is supported by the line shape of spectra around $(0, -0.6\pi)$, where the intensity at the Fermi level peaks along ΓX (see Fig. 3, inset). The small but distinct Fermi step is inconsistent with the formation of a pseudogap driving the dichotomy of the spectral weight as

reported for doped Sr_2IrO_4 [14]. Instead, the reduced weight along ΓX can be understood as the tail of a pole in the spectral function above the chemical potential arising from the merging of electron pockets from different M points, as seen in the LDA + SO + U calculations for $x \approx 0.15$. Hence, we conclude that La doped $\text{Sr}_3\text{Ir}_2\text{O}_7$ shows closed electronlike Fermi pockets with a double-lens shape.

The Fermi surface pockets are significantly smaller than half of the Brillouin zone, rendering the fermiology of doped $\text{Sr}_3\text{Ir}_2\text{O}_7$ qualitatively different from cuprates, which show open Fermi arcs at low doping, evolving into a large Fermi surface of volume $(1+x)$ near optimal doping [17,18].

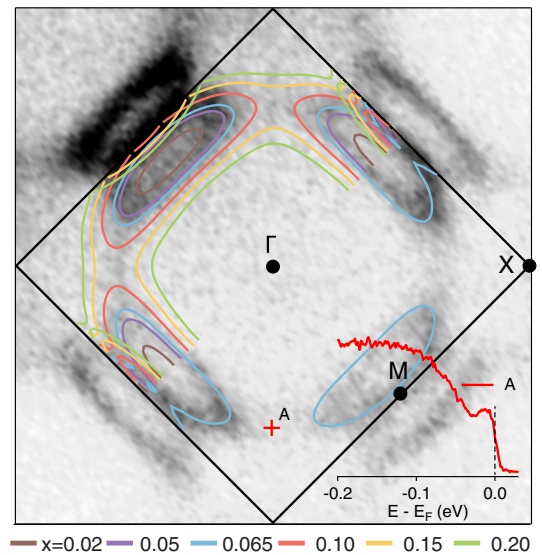


FIG. 3 (color online). LDA + SO + U calculations of the Fermi surface for $U = 3$ eV and different doping levels treated by the VCA superimposed on ARPES data for $x = 0.065$. The inset shows an EDC taken at 8 K at $\approx(0, -0.6\pi)$, demonstrating the absence of a pseudogap in the antinodal direction at the measurement temperature of 8 K.

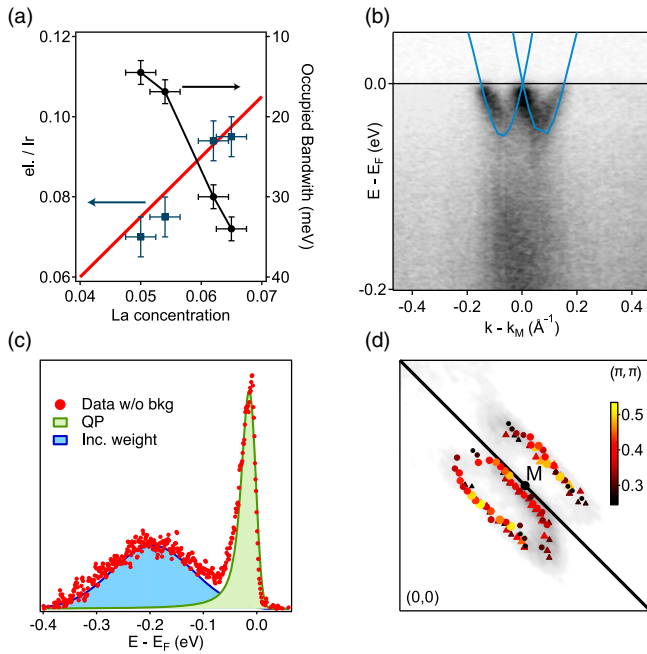


FIG. 4 (color online). Luttinger volume and quasiparticle residue. (a) Luttinger volume of the experimental Fermi surface (blue), compared with the nominal Luttinger volume of $3x/2$ arising from the stoichiometry of $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ if every lanthanum atom dopes one itinerant electron (red line). The doping evolution of the conduction band minimum is shown in black on the right axis. (b) Band dispersion along ΓM together with a LDA + SO + VCA calculation ($U = 0$, $x = 0.065$) along $(0, 0, -2\pi/c) - (2\pi/a, 0, -2\pi/c)$ corresponding to the ΓM direction of the 2D surface Brillouin zone. The hole pocket found in this calculation (see the Supplemental Material [38]) has been suppressed for clarity and the electron pocket at M was shifted in energy to match the experimental Fermi wave vectors. (c), (d) Quasiparticle residue Z along the Fermi surfaces at $\pm(\pi/2, \pi/2)$ (circles and triangles, respectively) estimated from the weight of the coherent quasiparticles.

We have extracted the Luttinger volume in $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ from extensive fits to the experimental Fermi surface. As shown in Fig. 4(a), the values are fully consistent with a small Fermi surface of volume $3x/2$, where x is the independently measured La concentration. The factor of $3/2$ arises from the stoichiometry of $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ and indicates that every La ion contributes one electron to the Fermi sea. The interpretation of our data within a conventional Fermi liquid picture with small Fermi surfaces is supported by a calculation of the electronic specific heat from the ARPES data. Using the experimental Fermi surface volume A_{FS} and quasiparticle velocities for $x = 0.065$ we calculate cyclotron masses $m^* = \hbar^2 \partial A_{\text{FS}} / 2\pi \partial \epsilon = 3.1(9) m_e$ for each Fermi surface lens. The doping dependence of m^* is negligible within the current accuracy of our experiment. Assuming two dimensionality, these masses correspond to a Sommerfeld coefficient $\gamma^{\text{ARPES}} = 9(3) \text{ mJ/mol K}^2$, in good agreement with the direct thermodynamic

measurement of $\gamma = 8.4\text{--}11.5 \text{ mJ/mol K}^2$ for different metallic samples, respectively [32]. This demonstrates that all low-energy charge excitations arise from the small Fermi surface pockets identified above. Furthermore, it strongly suggests that our ARPES data represent the bulk electronic structure.

The role of correlations in the low-energy excitations can be estimated from the renormalization of the Fermi velocity v_F/v_{bare} and the quasiparticle weight Z . While v_{bare} cannot be determined directly from experiment, it can often be approximated by band structure calculations within the LDA. In Fig. 4(b) we thus compare the experimental quasiparticle dispersion for $x = 0.065$ with a LDA + SO + VCA calculation for the same doping and relaxed atomic positions. Using averaged Fermi velocities from all Fermi surface crossings we find $v_F/v_{\text{bare}} = 0.5(2)$. The error bar is estimated from a slight uncertainty in the experimental Fermi velocity due to finite resolution effects and the high sensitivity of the calculated dispersion to the rotation angle of the octahedra. We further estimate the quasiparticle residue Z directly by analyzing the coherent weight. To this end we subtract a smooth background from the raw EDCs and fit the resulting spectra along the entire Fermi surface with two peaks representing the coherent quasiparticle and incoherent hump [see Fig. 4(c)]. While this analysis is somewhat model dependent and tends to underestimate Z , it clearly shows a significant coherent weight $Z = 0.25\text{--}0.5$ along the entire Fermi surface. This indicates a Fermi-liquid-like state with $Z \approx v_F/v_{\text{bare}}$ and thus a weakly momentum dependent self-energy, in stark contrast to lightly doped cuprates where strongly momentum dependent interaction effects lead to $Z \ll v_F/v_{\text{bare}}$ for the nodal quasiparticles [17,36].

The role of correlations in $\text{Sr}_3\text{Ir}_2\text{O}_7$ is thus rather intricate. On one hand, they are crucial for driving the insulating ground state in the parent compound via a substantial orbital dependent shift and deformation of the bare band structure [10,11]. On the other hand, they appear to play a minor role in the low-energy physics once metallicity is induced by light electron doping with La.

In summary, we investigated the doping evolution of the correlated insulator $\text{Sr}_3\text{Ir}_2\text{O}_7$. Our ARPES data show coherent, ungapped quasiparticles with a large and nearly momentum independent spectral weight forming a closed small Fermi surface, which can be accounted for by LDA + SO + U calculations. This phenomenology is in stark contrast to lightly doped cuprates [16–18] and surface doped Sr_2IrO_4 [14], suggesting that the properties of $\text{Sr}_3\text{Ir}_2\text{O}_7$ are unique among doped correlated insulators and not suitable for the engineering of cupratelike high- T_c superconductivity. It remains an open question how our findings relate to recent reports of strongly correlated states in doped single layer iridates [14,15] and Ru-substituted bilayer iridates [19].

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