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A unifying phase diagram with correlation-driven superconductor-to-insulator transition for the 122* series of iron-chalcogenides

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The 122* series of iron-chalcogenide superconductors, for example $K_xFe_{2-y}Se_2$, only possesses electron Fermi pockets. Their distinctive electronic structure challenges the picture built upon iron pnictide superconductors, where both electron and hole Fermi pockets coexist. However, partly due to the intrinsic phase separation in this family of compounds, many aspects of their behavior remain elusive. In particular, the evolution of the 122* series of iron-chalcogenides with chemical substitution still lacks a microscopic and unified interpretation. Using angle-resolved photoemission spectroscopy, we studied a major fraction of 122* iron-chalcogenides, including the isovalently 'doped' $K_xFe_{2-y}Se_{2-z}S_z$, $Rb_xFe_{2-y}Se_{2-z}Te_z$ and $(Ti,K)_xFe_{2-y}Se_{2-z}S_z$. We found that the bandwidths of the low energy Fe $3d$ bands in these materials depend on doping; and more crucially, as the bandwidth decreases, the ground state evolves from a metal to a superconductor, and eventually to an insulator, yet the Fermi surface in the metallic phases is unaffected by the isovalent dopants. Moreover, the correlation-driven insulator found here with small band filling may be a novel insulating phase. Our study shows that almost all the known 122*-series iron chalcogenides can be understood *via* one unifying phase diagram which implies that moderate correlation strength is beneficial for the superconductivity.

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I. INTRODUCTION

The iron-based superconductors (FeHTSs) can essentially be classified into two large families based on their electronic structure. The first family is comprised of iron pnictides and Fe(Te,Se) bulk crystals, whose Fermi surfaces are composed of both electron and hole pockets¹⁻³. Built on the spin fluctuations between the electron and hole Fermi pockets, a sign-changing s -wave pairing was predicted by weak coupling theories⁴⁻⁶. The second family consists of $A_xFe_{2-y}Se_2$ ($A=K, Rb, Cs, Ti/K$)⁷⁻¹⁰, $(Li_{0.8}Fe_{0.2})OHFeSe$ ^{11,12}, the ammonia-intercalated $Li_x(NH_2)_y(NH_3)_{1-y}Fe_2Se_2$ ¹³, and the recently discovered monolayer FeSe thin film¹⁴⁻¹⁷. The Fermi surfaces of this family of FeHTSs consist only of electron pockets. The nodeless gap observed in these iron-chalcogenides has posed a serious challenge to the weak coupling theories, which have predicted gap nodes for such a Fermi surface topology¹⁸. Therefore, a comprehensive understanding of microscopic behaviors of both families is critical for understanding the mechanism of FeHTS.

Recently, a systematic angle-resolved photoemission spectroscopy (ARPES) investigation was conducted in many series of the first families of FeHTSs, including bulk Fe(Te,Se), $BaFe_2(As_{1-x}P_x)_2$, $Ba(Fe_{1-x}Ru_x)_2As_2$, $NaFe_{1-x}Co_xAs$ and $LiFe_{1-x}Co_xAs$ ¹⁹. In all these compounds, the bandwidths are found to increase with the doping level x , as the superconductivity is weakened and eventually suppressed at high dop-

ing levels, regardless of the doping traits being heterovalent or isovalent. Such an observation is consistent with strong coupling theories, where pairing is mediated by local antiferromagnetic interactions²⁰⁻²⁶. It is then intriguing to ask if similar behavior is also present for the second family of FeHTS. However, in contrast to the first family of FeHTS, an in-depth understanding of the doping dependence of the second family of FeHTS is still lacking for several reasons. The doping of many members of the second family is discrete or cannot be controlled. In particular, for the 122* series of iron-chalcogenides ($A_xFe_{2-y}Se_2$ and the like) that comprise most of this family, there is an intrinsic phase separation between a $\sqrt{5} \times \sqrt{5}$ Fe-vacancy ordered insulating phase (also called the "245" phase) and a superconducting phase²⁷⁻³¹. Consequently, the doping-dependence of the microscopic properties of this series has not been investigated extensively.

Recent progress in materials synthesis has enabled the tuning of the superconductivity of $A_xFe_{2-y}Se_2$ in a more controlled manner. As shown in Fig. 1(a), phase diagrams have been obtained when replacing Se with Te or S in $A_xFe_{2-y}Se_2$ ($A = K, Rb$)^{32,33}. Intriguingly, though the chemical pressure is tensile for Te doping and compressional for S doping as shown by their lattice constant evolution in Fig. 1(b), superconductivity is gradually suppressed in both cases. See the Supplementary Material for in-plane resistivity and magnetic susceptibility data³⁴. It is important to note that at low temperature the resistivity on the S-overdoped side is metallic, while

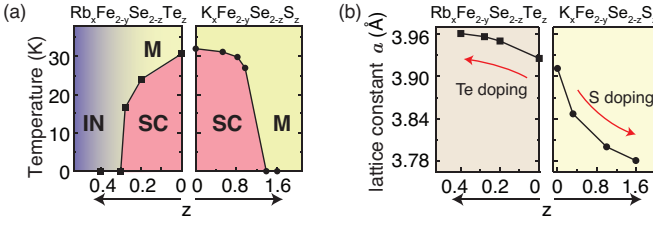


FIG. 1: (a) Phase diagrams of $\text{Rb}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{Te}_z$ and $\text{K}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{S}_z$. SC, M, and I represent the superconducting, metallic and insulating phases, respectively. (b) In-plane lattice constants (data are extracted from Ref.^{32,33}) of $\text{Rb}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{Te}_z$ and $\text{K}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{S}_z$.

that on the Te-overdoped side is insulating.

In this work, we systematically studied the low energy electronic structure of $A_x\text{Fe}_{2-y}(\text{Se}, \text{S/Te})_2$ ($A=\text{K}, \text{Rb}, \text{Ti/K}$), and we show that they can be fit into one generic phase diagram, with bandwidth or correlation strength being the control parameter. We found that on decreasing bandwidth, the ground state of these compounds evolves from a metal to a superconductor and then to an insulator, which has almost all the essential ingredients of the cuprate phase diagram, except that bandwidth, rather than carrier doping, is the control parameter. Intriguingly, the insulating phase found here has a small band filling and is induced by band narrowing from a superconducting phase, which resembles the bandwidth-controlled Mott transition, except that the latter requires an *integer* band filling. Therefore it might indicate either a novel correlation-driven insulator, or that the unit cell has been somehow expanded by certain ordering. Our results give a microscopic and unified understanding of various chemical-substitution-based phase diagrams of the 122* series of iron-chalcogenide FeHTS. The revealed relationship between the superconductivity and the electronic correlations is reminiscent of similar observations made in iron pnictides¹⁹, and thus might lead to a comprehensive understanding of both families of FeHTS.

II. II. EXPERIMENT

High-quality $\text{K}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{S}_z$ ($z = 0, 0.53, 0.83, 0.97, 1.40$, and 1.61) single crystals were synthesized using the self-flux method. High-purity Fe, Se, S and K were carefully mixed with the nominal compositions, then sealed in an evacuated quartz tube. The tube was heated to 1303 K and kept for 6 h, and then slowly cooled down to 1003 K at the rate of 3 K/h before shutting off the power. The S concentrations, z , were confirmed by electron-probe micro-analysis (EPMA). Further details about the $\text{Rb}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{Te}_z$ ($z = 0, 0.20, 0.28, 0.3$, and 0.4) and $\text{Ti}_{0.4}\text{K}_{0.4}\text{Fe}_{1.7}\text{Se}_2$ single crystals can be found in Refs.^{32,35}. $\text{Ti}_x\text{Fe}_{2-y}\text{Se}_{0.4}\text{S}_{1.6}$ and $\text{Ti}_x\text{Fe}_{2-y}\text{S}_2$ are non-superconducting compounds³⁶. High-resolution ARPES measurements were performed at the I05 beamline of the Diamond Light Source, beamline 5-4 of the Stanford Synchrotron Radiation Light source (SSRL), One-Cubed ARPES at BESSY II and our in-house ARPES system with a Helium discharge lamp (21.2 eV photons), all equipped with a Scienta R4000

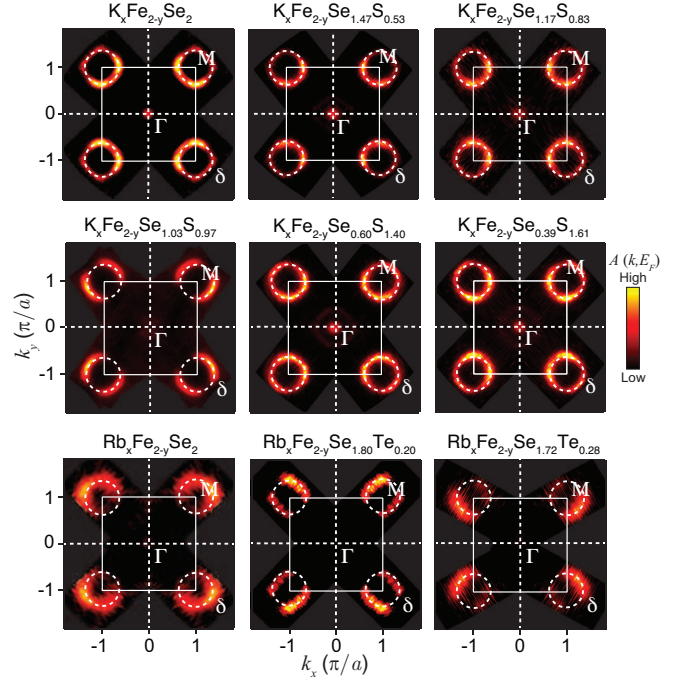


FIG. 2: Doping dependence of the Fermi surface sheets of Te/S doped $A_x\text{Fe}_{2-y}\text{Se}_2$. These nine false-color plots are the four-fold-symmetrized photoemission intensity maps at the Fermi energy (E_F) in the Γ - M plane. The photoemission intensity distribution is mirror symmetrized with respect to the k_x and k_y axes. Intensities were integrated over a window of $(E_F - 15 \text{ meV}, E_F + 15 \text{ meV})$. White dashed circles show the δ band Fermi crossings, wherever identifiable.

electron analyzer, and at beamline 28A of the Photon Factory, KEK, equipped with a Scienta SES-2002 electron analyzer. The overall energy resolution was 2.5~20 meV, depending on the photon energies and experimental setups, and the angular resolution was 0.3 degrees. Samples were cleaved under ultra-high vacuum conditions. $\text{Rb}_x\text{Fe}_{2-y}\text{Se}_{1.6}\text{Te}_{0.4}$ samples are measured at $T=100 \text{ K}$ to avoid charging effects and other samples are measured at $T = 1\sim 13 \text{ K}$ depending on the experimental setups. The ARPES measurements on each sample were carried out within 8 hours, with sample aging effects carefully monitored.

III. III. RESULTS

We first examine the Fermi surface topologies of $\text{K}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{S}_z$ and $\text{Rb}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{Te}_z$ in Fig. 2. Similar to $\text{K}_x\text{Fe}_{2-y}\text{Se}_2$ ⁷⁻⁹, only electron pockets can be observed at the zone center Γ and the zone corner M . There are two electron-like bands δ and δ' around the M point, which give nearly the same Fermi surface sheets^{37,38}. However, only the shallower δ band can be clearly resolved [see Figure. 3(a)] in our experimental geometry as reported before⁷. Around the zone center Γ , the shallow electron-like band κ contributes a small amount of photoemission intensity, with a strong k_z dispersion as observed before in $\text{K}_x\text{Fe}_{2-y}\text{Se}_2$ ⁷. We present the k_z dispersions

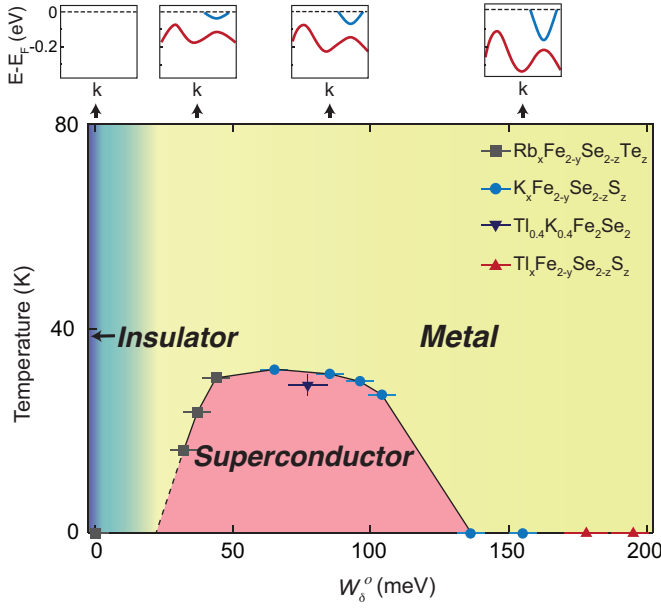


FIG. 4: A unified phase diagram of T_c vs. W_δ^0 for $A_x\text{Fe}_{2-y}(\text{Se,S})\text{Te}_2$ ($A=\text{K, Rb, Tl/K}$). The sketches of the band dispersions for several dopings illustrate the band structure evolution and the absence of quasiparticles in the insulator phase. The dashed line is a linear extrapolation of the T_c vs. W_δ^0 data. Note that the bandwidths for $(\text{Tl,K})_x\text{Fe}_{2-y}(\text{Se,S})_2$ are extrapolated to match the doping level of the other materials³⁴.

along the Γ - M direction. Here, the bandwidth of the β band can be clearly resolved, and increases monotonically from left to right, in conjunction with the reduction of the in-plane lattice constant. The bottom of the δ band sinks accordingly. Such a band structure evolution is further illustrated by energy distribution curves (EDCs) in Fig. 3(b), where a background has been subtracted to retrieve the dispersion more clearly as described in the Supplementary Material³⁴. Figure 3(c) summarizes the systematic evolution of the bandwidth of β (W_β) and the occupied width of the δ band (W_δ^0) with S/Te doping, where W_β is enhanced by 120 %, and W_δ^0 by a factor of more than 4, on decreasing the lattice constants or correspondingly, increasing the chemical pressure.

Figure 3(d) shows the photoemission spectra along Γ - M for $\text{Rb}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{Te}_z$ ($z=0.3, 0.4$). The spectra do not show any noticeable momentum dependence. There are no quasiparticles around the chemical potential, as expected for insulators. However, the $z=0.3$ sample has some residual spectral weight near the chemical potential, while the $z=0.4$ sample has none [Fig. 3(e)]. This may due to the coexistence of some trace amount of superconducting phase in the $z=0.3$ sample as indicated in the resistivity data³², or the incoherent remains of the suppressed quasiparticle bands. Moreover, our high-resolution transmission electron microscope (HRTEM) data of $\text{Rb}_x\text{Fe}_{2-y}\text{Se}_{1.6}\text{Te}_{0.4}$ show that the $\sqrt{5} \times \sqrt{5}$ vacancy-ordered “245” insulating phase coexists with a $\sqrt{2} \times \sqrt{2}$ superstructured phase in mesoscopic scale³⁴. Since a similar $\sqrt{2} \times \sqrt{2}$ superstructure exists in the superconducting phase, as shown by the band folding in Fig. 3(a), one can conclude that this

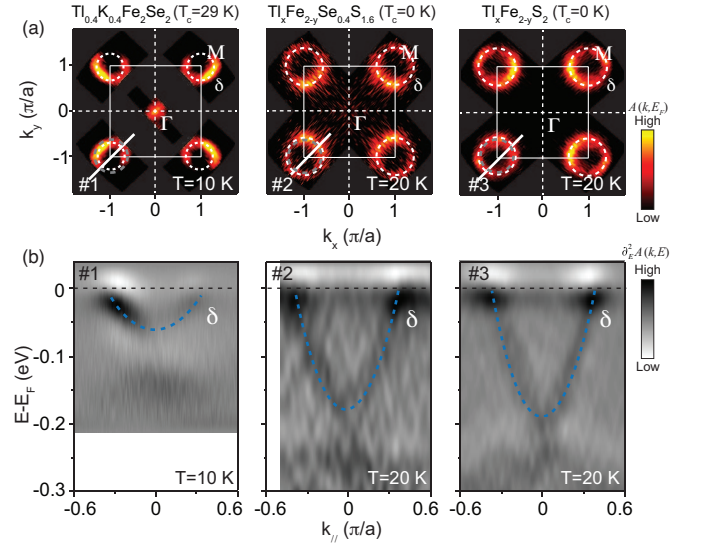


FIG. 5: ARPES data on $\text{Tl}_{0.4}\text{K}_{0.4}\text{Fe}_{1.7}\text{Se}_2$, $\text{Tl}_x\text{Fe}_{2-y}\text{Se}_{0.4}\text{S}_{1.6}$ and $\text{Tl}_x\text{Fe}_{2-y}\text{S}_2$. (a) False-color plots of the four-fold symmetrized photoemission intensity maps. Intensities were integrated over a window of $(E_F - 15 \text{ meV}, E_F + 15 \text{ meV})$. White dashed circles show the δ Fermi crossings, while the electron pockets of $\text{K}_x\text{Fe}_{2-y}\text{Se}_2$ are illustrated as dashed blue circles. (b) Photoemission intensity distributions at the zone corners along the cuts illustrated by white solid lines in (a).

phase with the $\sqrt{2} \times \sqrt{2}$ superstructure evolves from a superconductor to an insulator with increasing Te doping. This insulator is distinct from that induced by Fe vacancies in the “245” phase^{29,43,44}. Moreover, since Te only introduces chemical pressure instead of altering the band filling, such a sudden disappearance of Fermi surface when the bandwidth diminishes indicate a correlation-driven superconductor/metal-insulator transition. Due to the strong competitions among various inter-site exchange interactions and strong spin fluctuations in the iron chalcogenides, as exemplified in bulk FeSe ^{45,46}, it is an open question whether there is a magnetic order in this insulator phase with the $\sqrt{2} \times \sqrt{2}$ superstructure. This might be answered by future neutron scattering studies.

As shown in Ref. ³⁴, band calculations indicate that the bare bandwidths of KFe_2Te_2 , KFe_2Se_2 and KFe_2S_2 only differ by about 63% (W_β) and 19% (W_δ^0)⁴¹. By applying a linear interpolation, we estimate that the change of the bare bandwidth in the measured doping range here should be within 32% (W_β) and 10% (W_δ^0). The remarkable change of the bandwidths in Fig. 3 is thus due to the additional bandwidth renormalization by electronic correlations, which is driven largely by Hund’s rule coupling in the FeHTSs⁴⁷. Similar behaviors have been observed in the prototypical bandwidth-controlled Mott transition of $\text{NiS}_{2-x}\text{Se}_x$ ⁴⁸. Because the bare bandwidth and the effective bandwidth are monotonically related, either of them can be used to characterize the itinerancy or kinetic energy. However, since the effective bandwidth not only represents the itinerancy of quasiparticles, but also carries the information on the electronic correlations in these compounds, we use the effective bandwidth in the following discussions.

In addition to the bandwidth, effective mass (m^*) and

Fermi velocity (v_F) of the δ band reflect the electronic correlation strength as well. As shown in Fig. 3(f), m^* decreases from the Te-overdoped side to the S-overdoped side, while v_F shows the opposite trend accordingly. Taking W_δ^O as the electronic parameter, one can obtain the phase diagram in Fig. 4, which shows that superconductivity is suppressed by driving the bandwidth away from its optimal value by doping either the isovalent S or Te. Furthermore, we have examined whether other sibling iron-chalcogenides can fit into this phase diagram. Data for $\text{Ti}_{0.4}\text{K}_{0.4}\text{Fe}_{1.7}\text{Se}_2$ ($T_c = 29$ K) and $\text{Ti}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{S}_z$ (non-superconducting) are shown in Figs. 5(a) and (b). Compared with $\text{K}_x\text{Fe}_{2-y}\text{Se}_2$, the electron Fermi pockets of $\text{Ti}_{0.4}\text{K}_{0.4}\text{Fe}_{1.7}\text{Se}_2$ are smaller [Fig. 5(a)], corresponding to a carrier concentration of 0.14 ± 0.02 electrons per Fe. However, assuming these off-plane dopants do not affect the band renormalization, as found for $\text{Ba}_{1-x}\text{K}_x\text{Fe}_2\text{As}_2$ before, we rigidly shift the parabolic band so that the extrapolated Fermi crossings match those of $\text{K}_x\text{Fe}_{2-y}\text{Se}_2$ ³⁴, the extrapolated W_δ^O would fall into the curve shown in Fig. 4 within the error bar. For the non-superconducting $\text{Ti}_x\text{Fe}_{2-y}\text{Se}_{0.4}\text{S}_{1.6}$ and $\text{Ti}_x\text{Fe}_{2-y}\text{Se}_2$, their electron pockets are slightly larger than those of $\text{K}_x\text{Fe}_{2-y}\text{Se}_2$, while their δ bands are much broader (Fig. 5). After compensating the small difference in the Fermi pockets by rigidly shifting their Fermi crossings to match those of $\text{K}_x\text{Fe}_{2-y}\text{Se}_2$, one can obtain their W_δ^O s. They are still much larger than 150 meV, far above the superconducting regime in Fig. 4. We note that a similar phase diagram can be reached if using m^* as the horizontal axis.

IV. IV. DISCUSSIONS

We have observed the bandwidth evolution of the 122* series of iron-chalcogenides by substituting Se with S or Te, and obtain a unified phase diagram in Fig. 4 from metal (S side) to superconductor, then to an insulator (Te side), with nearly fixed Fermi surface and negligible impurity scattering effects (see Supplementary Materials³⁴). Moreover, there is no noticeable orbital occupation change either. The dramatic change of bandwidth (or band bottom of δ) signifies strong correlation effects in iron-selenides. Such a bandwidth-controlled or correlation-driven superconductor-to-insulator transition resembles the bandwidth-controlled or the Brinkman-Rice Mott transition, where the bandwidth decreases to zero or the quasiparticle mass diverges⁴⁹. Interestingly, a Mott transition, no matter in a single-band or multiple-band system, requires integer filling of the band^{49,50}, whereas there is a small band filling in the insulating state of Te-overdoped sample. Therefore, it is not a Mott insulator or at least not a usual one, instead, it could be a novel correlation-driven insulator where Anderson localization might play a role. Alternatively, judging from the depleted spectral weight at the chemical potential, it is also likely that certain order, such as charge density wave, might have occurred and multiplied the unit cell, which in turn would cause an integer band filling, and give a Mott transition. Recently, J. He and coauthors reported a carrier-doping-induced insulating phase

in monolayer FeSe ⁵¹. However, it was later found by Y. Fang and coauthors that similar band structure and insulating phase could be reproduced in FeSe films with iron vacancies⁵². Therefore, the insulating phase in FeSe film is qualitatively different from the insulating $\text{Rb}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{Te}_z$ ($z=0.3, 0.4$).

In Fig. 4, there is a metallic regime between the insulating phase and superconducting phase. If this regime existed in real materials, it would correspond to the narrow tellurium concentration range of $0.28 < z < 0.3$. Since this doping range is quite narrow, the sudden change between insulating state and superconducting state indicates that the superconductor-to-insulator transition here is likely of first order, and this metallic regime may not exist in real materials. Consistently in this narrow doping range, the transport behaviors evolve drastically, and there is some trace of the superconducting phase, an indicative of phase separation³².

Towards the large bandwidth or weak correlation regime, the behavior of the 122* series of iron-chalcogenides resembles the observations made before in the first family of FeHTS¹⁹. That is, the superconductivity is weakened and eventually suppressed at the high doping level with increased bandwidth. However, in the strong correlation regime, most iron pnictides enters the competing collinear antiferromagnetic metallic ground state instead of an insulating state found here. Nevertheless, all these results jointly suggest that the bandwidth or correlation strength provides a unifying standpoint in understanding the ground-state evolutions in both families of FeHTSs.

Like the t - J model for the cuprate, FeHTSs have been described by effective models with short-range antiferromagnetic exchange couplings in strong local pairing scenarios, which could give the pairing symmetries and gap functions for both families of FeHTSs²⁰⁻²⁶. In such effective models, there are two critical parameters that control the superconductivity: the renormalized effective bandwidth, W , and the effective antiferromagnetic coupling strength J (or multiple J_i s among various neighbors). The superconductivity appears when the ratio of J/W is bound within a certain region. Below the lower bound, electron correlations are too weak, so that the system cannot sustain the unconventional high temperature superconductivity; while beyond the upper bound, too strong electron correlations or too weak itinerancy would give an insulator and/or magnetic ordered phase instead of a superconductor. If the effective short-range antiferromagnetic exchange couplings J s do not vary significantly with the isovalent doping, as in the case of FeHTSs^{53,54}, the bandwidth (or other equivalent parameters describing electron itinerancy, or conversely, correlation) becomes the pivotal variable that controls the evolution of superconductivity, as observed here in these 122* series of iron-chalcogenide superconductors. Therefore, the observed robust correlation between the superconductivity and W provides a compelling support for the local pairing scenario in the 122* series of iron-chalcogenides superconductors. Furthermore, in the strong coupling theories that tried to understand both FeHTSs and cuprates^{22,26}, the phase evolution is controlled by the competition between the kinetic energy and interactions. Therefore, it is not surprising that the phase diagram found here resembles the cuprate phase diagram, except

that the control parameter here is bandwidth, instead of carrier doping. Note that in the cuprate case, with increased doping, the effective bandwidth increases, and the correlation strength decreases. As a result, it is possible to understand both the FeHTSs and the cuprates in such a unified way.

Finally, we note that our data further challenges the weak coupling mechanism, which stresses the interactions of electrons around the Fermi surface. In this kind of scenarios, the stronger correlations (or narrower bandwidth, smaller v_F) give rise to a larger DOS, which is expected to yield higher T_c , when the Fermi surface is invariant upon isovalent doping. However, our data suggests that the superconductivity will be strongly suppressed once the electron correlation strength goes beyond the optimal regime, even though the DOS continuously increases. For example, $\text{Rb}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{Te}_z$ has stronger correlations but a lower T_c than $\text{Rb}_x\text{Fe}_{2-y}\text{Se}_2$. Moreover, $\text{Ti}_{0.4}\text{K}_{0.4}\text{Fe}_{1.7}\text{Se}_2$ and $\text{K}_x\text{Fe}_{2-y}\text{Se}_2$ have almost the same T_c , m^* and v_F [dashed line in Fig. 3(f)], while there is significant difference between their sizes of Fermi surfaces. This is not consistent with the weak coupling mechanism as well.

V. V. CONCLUSIONS

To conclude, we have shown that the superconductivity evolutions in $\text{K}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{S}_z$, $\text{Rb}_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{Te}_z$ and $(\text{Ti,K})_x\text{Fe}_{2-y}\text{Se}_{2-z}\text{S}_z$ can be comprehensively understood from

their bandwidth evolutions. We have found a correlation driven phase evolution from metal to superconductor, and eventually to a non-trivial insulating phase. Our data suggest that correlation (or bandwidth as the electronic control parameter) plays a critical role in the iron based superconductors with only electron Fermi surfaces, as found before for those with both electron and hole Fermi surfaces. Our results thus present an important step to the unified understanding of all the iron-based superconductors, which are consistent with the strong local pairing scenarios.

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