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# Anomalous doping evolution of nodal dispersion revealed by *in-situ* ARPES on continuously doped cuprates

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## Abstract

We study the systematic doping evolution of nodal dispersions by *in-situ* angle-resolved photoemission spectroscopy on the continuously doped surface of a high-temperature superconductor  $\text{Bi}_2\text{Sr}_2\text{CaCu}_2\text{O}_{8+x}$  and reveal that the nodal dispersion has three fundamentally different segments separated by two kinks, located at  $\sim 10$  meV and roughly 70 meV, respectively. These three segments have different band velocities and different doping dependence. In particular, in the underdoped region the velocity of the high-energy segment increases monotonically as the doping level decreases and can even surpass the bare band velocity. We propose that electron fractionalization is a possible cause for this anomalous nodal dispersion and may even play a key role in the understanding of exotic properties of cuprates.

For the high-temperature (high- $T_c$ ) cuprates, one of the puzzling features is the “kink” in the electronic band dispersion, which universally exists in different superconducting families of cuprates [1-3]. Along the nodal direction of the d-wave superconducting gap, the kink appears at a binding energy of roughly 70 meV in many cuprates [1,4-6]. Meanwhile near the antinode with the maximum energy gap, a seemingly stronger kink is located at about 20-40 meV [7-9], depending on the doped carrier concentration  $x$ . Many scenarios have been proposed to understand the origin of this kink feature [10-12]. For examples, one popular proposal is that a strong electron-phonon coupling induces a kink in the dispersion [13,14], while another one suggests that the electron-magnetic mode coupling is responsible for the kink [15-17]. Yet, the origin of this universal kink is far from settled. Recently we developed a new technique to continuously change the doping level of surface layers by annealing sample in ozone/vacuum atmosphere, which enables a systematic *in-situ* angle-resolved photoemission spectroscopy (ARPES) study on the same surface with a wide range of doping level, thus promising more precise studies on doping evolution of many important properties of the cuprates [18].

In this letter, we systematically study the doping evolution of the nodal dispersion by performing *in-situ* ARPES measurements on a continuously doped surface of  $\text{Bi}_2\text{Sr}_2\text{CaCu}_2\text{O}_{8+x}$  (Bi2212). From the measured band dispersions, two kinks are identified at  $\sim 10$  meV and 45-70 meV below the Fermi level ( $E_F$ ), respectively, and they separate the nodal dispersion into three segments with different band velocities. We clearly demonstrate the fundamentally different doping evolutions for these three band velocities: a decreasing velocity for the lower segment, a no-change velocity for the middle segment, and an increasing velocity for the higher segment, as the doping level decreases. What is more exotic is that the velocity of the higher segment becomes divergent and can even exceed the bare band velocity for the extremely underdoped (UD) ones. After discussing several possible scenarios proposed to explain the kink phenomena, we suggest that electron fractionalization at the high energy is likely responsible for the anomalous doping evolution of the nodal dispersion in the UD region.

Sample preparation and surface treatment of ozone/vacuum annealing to continuously change the doping level of surface layers were described in our previous paper [18]. *In-situ* ARPES measurements were performed in a laboratory ARPES system equipped with a Scienta R4000 analyzer and a Scienta VUV light source. He-I $\alpha$  resonant line ( $h\nu = 21.218$  eV)

was used and the vacuum of the ARPES chamber was better than  $3 \times 10^{-11}$  Torr. The energy and angle resolution were set as  $\sim 5$  meV and  $\sim 0.2^\circ$ , respectively. All data in this paper were acquired at 12K to make sure the sample surface at all doping levels is in the superconducting state.

We first show the spectrum image plots with the same color scale along the nodal direction at various different doping levels from  $x \sim 0.24$  to 0.08 [Figs. 1(a)-1(f)], which were acquired on the same surface through annealing sample in the ozone/vacuum circumstance [19]. The well-known 70-meV kink is clearly observed at each doping level, and the spectral intensity has a sudden reduction after crossing the kink. The area with high intensity (light color in images) shrinks dramatically when it goes to the UD region. Interestingly, the band beyond the kink becomes more and more vertical, which indicates that the kink becomes stronger with decreasing doping level. In order to extract the band dispersion for quantitative comparison, a standard practice [20] is to perform a Lorentzian fitting to momentum distribution curves (MDCs) that are intensity distributions as a function of momentum at a fixed energy [Figs. 1(g)-1(i)], with an assumption that the self-energy (SE) in Green's function has no or weak dependence on momentum. From the fitting, the peak positions of MDCs trace out the band dispersion and the widths contain the information of the quasiparticle lifetime. The extracted band dispersion for each doping level is plotted with the black lines in Figs. 1(a)-1(f). Again, one can clearly see that the kink phenomena become stronger with decreasing doping level. The spectral width is also seen to become wider as the doping level decreases [Figs. 1(g)-1(i)]. Meanwhile, the spectral width becomes larger suddenly when it passes across the kink [Figs. 1(j)-1(k)], indicating an abrupt change of the quasiparticle lifetime.

In principle, ARPES probes the single-particle spectral function, which treats the many-body interactions as additional SE renormalization to electrons [21]. One way to understand the interactions in a material is to analyze the SE  $\Sigma(w)$  from ARPES spectra. By comparing the measured band with its corresponding bare band [Fig. 2(a)], one can extract the real part  $\text{Re}\Sigma$ , which is the energy difference between the measured band and the bare band, and the imaginary part  $\text{Im}\Sigma$ , which is proportional to the MDC width. According to previous practices [22], the low-energy bare band along the nodal direction in the cuprates can be approximated as a linear line that connects the measured Fermi crossing point and the higher binding energy, typically around  $\sim 200$  meV. We note that there are other different forms for the bare band used in

previous literature [23]. For direct comparison, the MDC-derived nodal dispersions at different doping levels are plotted with the relative momentum to their corresponding Fermi momenta ( $k_F$ ) [Figs. 2(a) and 3(a)]. We notice that previous literature [1,4-9] usually plot the real part  $\text{Re}\Sigma$  and the MDCs' width in the binding energy [Figs. 2(b) and 2(d)] due to the weak dependence on momentum. In order to better visualize the SE, we also plotted  $\text{Re}\Sigma$  and MDCs' width against the peak positions of the fitted MDCs [Figs. 2(c) and 2(e)]. One can see the conspicuous peaks in  $\text{Re}\Sigma$  curves [Figs. 2(b) and 2(c)], which indicates that there is a large renormalization of the electronic dispersion at where the kink occurs. The energy position of the kink [Fig. 2(f)] and relative momentum position of the kink [Fig. 2(g)] for different doping levels, can be identified through these conspicuous peaks. With decreasing doping level, the peak in  $\text{Re}\Sigma$  curves gradually grows and moves to the  $E_F$  or  $k_F$  and its height is enhanced due to the stronger kink. Meanwhile the width of MDCs dramatically increases after the kink for the UD samples, while it is much milder before the kink [Figs. 2(d) and 2(e)]. This behavior becomes more and more noteworthy as the doping level decreases, especially in the UD side. It resembles that electrons are undergoing a decay from a well-defined quasiparticle to something with an extremely short lifetime when they go cross the kink. What is the more impressive is that the kink gradually moves toward their corresponding  $k_F$  and  $E_F$  with decreasing doping level in the UD side, which directly suggests the area with a well-defined quasiparticle shrinks [Figs. 2(f) and 2(g)].

Since the quasiparticle lifetime experiences a large transformation after crossing the kink, we next study the effective mass of the band dispersion before and after the kink. The MDC-derived nodal band dispersions at different doping levels are summarized in Figs. 3(a)-3(c). Besides the kink at 45-70 meV, one can see another kink of the band dispersion at a lower binding energy  $\sim 10$  meV [19], which contributes a "shoulder" in the  $\text{Re}\Sigma$  curves [Fig. 2(b)] and becomes more pronounced at the UD levels [Fig. 3(a)]. These two kinks separate the nodal band dispersion into three segments with different band velocities or effective masses. To make the results more precise, we avoid the vicinity of these two kinks and evaluate velocities of three band segments at the binding energy of 0-5 meV ( $v_l$ ), 25-55 meV ( $v_m$ ) and 80-125 meV ( $v_h$ ) [Fig. 3(d)]. We note that the energy range to evaluate the middle-range velocity varies slightly for different doping levels in order to avoid the nearby low-energy kink. The numerical analysis of doping dependence on the velocities of three band segments is displayed in Fig. 3(e).

We find that the  $v_l$  just below  $E_F$  decreases smoothly with decreasing doping level [Fig. 3(b)], which is consistent with the theoretical expectation that stronger electron-electron interactions at UD samples shall make electrons heavier [24]. The  $v_m$  remains a constant [Fig. 3(c)] over a wide doping range, consistent with previous observations [25,26]. The exotic behavior is that the  $v_h$  at the higher binding energy dramatically increases with decreasing doping level in the UD region, in opposite with the expected trend. More surprisingly, at the heavily UD level the  $v_h$  can even surpass its bare band velocity which is defined by LDA calculations [27,28]. Those exotic behaviors of the velocity imply an unconventional origin of the kink in the nodal dispersion.

The popular scenario of kink origin attributes these phenomena in electronic dispersion as the renormalized effect to the electrons due to the electrons coupling with some kinds of bosonic mode at specific energies, like phonons or spin resonance [1,13-17]. Although some of these bosonic modes do exist in the cuprates [29-31], these coupling scenarios are unable to explain the doping dependence of the kink phenomena, especially in the aspect of the divergent  $v_h$ . The magnetic resonance mode scenario is unlikely since the kink in the nodal dispersion still robustly survives in the normal states where the magnetic resonance mode vanishes [1,31], so here we focus on discussing the electron-phonon coupling (EPC) scenario. Firstly, under the frame of the EPC, typically the bare bands can be regarded approximately as a linear line within a narrow energy window that connects the measured  $E_F$  and a higher binding energy just like the dashed line in Fig. 2(a). Thus, with decreasing doping level, due to the increasing  $v_h$ , the bare bands will become more and more vertical, which is directly incompatible with the simple rigid band shift picture of Bi2212 [32,33]. Secondly, according to the weak EPC scenario described by Migdal-Eliashberg picture [34], since the EPC domain should be restricted to a small energy range around the typical energy of phonon modes ( $\Omega$ ), so the dispersion well above  $\Omega$  tends to be recovered to the position of the noninteracting band. Thus, the band velocity of the segment at a higher binding energy than  $\Omega$  will remain relatively stable, and for the binding energy lower than  $\Omega$ , the corresponding velocity will change with the coupling strength, which is opposite to what was observed here. Thirdly, we noticed the strong EPC picture due to the formation of the polarons [35,36] is argued to be a proper explanation for the kinks in the undoped and lightly UD cuprates. However, a placid change of quasiparticle lifetime at the kink point and a more placid evolution of high-energy band velocity as a function of doping level inferred from polaronic scenario cannot capture our measurements here.

On the contrary, it is reasonable that the kink at the lower energy  $\sim 10$  meV stems from the EPC [26,37,38] due that it fits well with the conventional coupling frame. The segments after the lower energy kink recovers to the non-coupling dispersion which has a constant velocity, while distinguishably the segments before the lower energy kink have different velocities due to variable strength of coupling. In fact, there is strong evidence of the existence of the lower energy phonons in Bi2212 [39].

We noticed that Randeria et al [11] phenomenologically suggested the  $v_h$  are a consequence of the renormalization to the nodal quasiparticle weight  $Z_k$  through a formula  $v_h = v_m/Z_k$ . Due that the  $v_m$  is constant, the nodal quasiparticle weight  $Z_k$  should decrease quite rapidly with decreasing doping level, which is not consistent with our previous results that the nodal quasiparticle weight  $Z_k$  keeps relatively constant over a wide doping range [18].

The unconventional behavior of the  $v_h$  surpassing the predicted velocity of the bare band is reminiscent of the electron fractionalization in some of the 1D materials [40,41]. Similar scenarios of electron fractionalization have also been proposed for the 2D cuprates [42-45]. For the parent compound of a cuprate superconductor, a Mott insulator, the charge is gapped due to strong on-site repulsive energy [46], while the spin has a gap-less excitation [47], so the charge and spin degrees can be separated in the low-energy excited processes, namely, the electrons are fractionalized. With the holes doped into, the system gradually evolves into a high- $T_c$  superconductor, and this fractionalization effect becomes weaker but still leaves some signatures in the UD region. In fact, there are many theoretical models suggesting that the spin-charge separation is the key to the cuprates [46,48-50]. One of these theoretical models is the so-called phase string theory [50]. In a doped Mott insulator described by the simplest t-J model [46], the doped holes cannot propagate coherently due to the presence of a nontrivial sign structure stemming from the phase string sign structure, which is distinct from the conventional fermion sign structure [50-52]. In order to incorporate the charge propagation properly, the doped holes should be fractionalized at a higher energy. The phase string theory [53] predicts that a peculiar electron fractionalization happens in the doping region accompany with the pseudo-gap, which intrinsically leads to a two-component structure: 1. The band segment before the 70 meV kink would be the “protected” emergent quasiparticle with the bare-band Fermi velocity, which is independent with the doped concentration  $x$ ; 2. The segment after the kink acquires a larger band velocity due to the natural electronic fractionalization, which increases

conspicuously with decreasing doping level, especially for very low doped ones. Naturally, due to the electron fractionalization, the spectra have a sudden broadening after crossing the kink. A comprehensive theoretical demonstration along this line is given in a separated paper [54], which nicely reproduced many observed features in our ARPES measurements.

In conclusion, we conduct a systematic ARPES study of nodal dispersion of Bi2212 over a wide doping range and reveal that the velocity of the segment after the high-energy kink unconventionally increases with decreasing doping level in the UD region, which cannot be explained by the electron-mode coupling scenario. Alternatively, we propose the electronic fractionalization effect at the high energy might be responsible for the unconventional behaviors.



Figures:

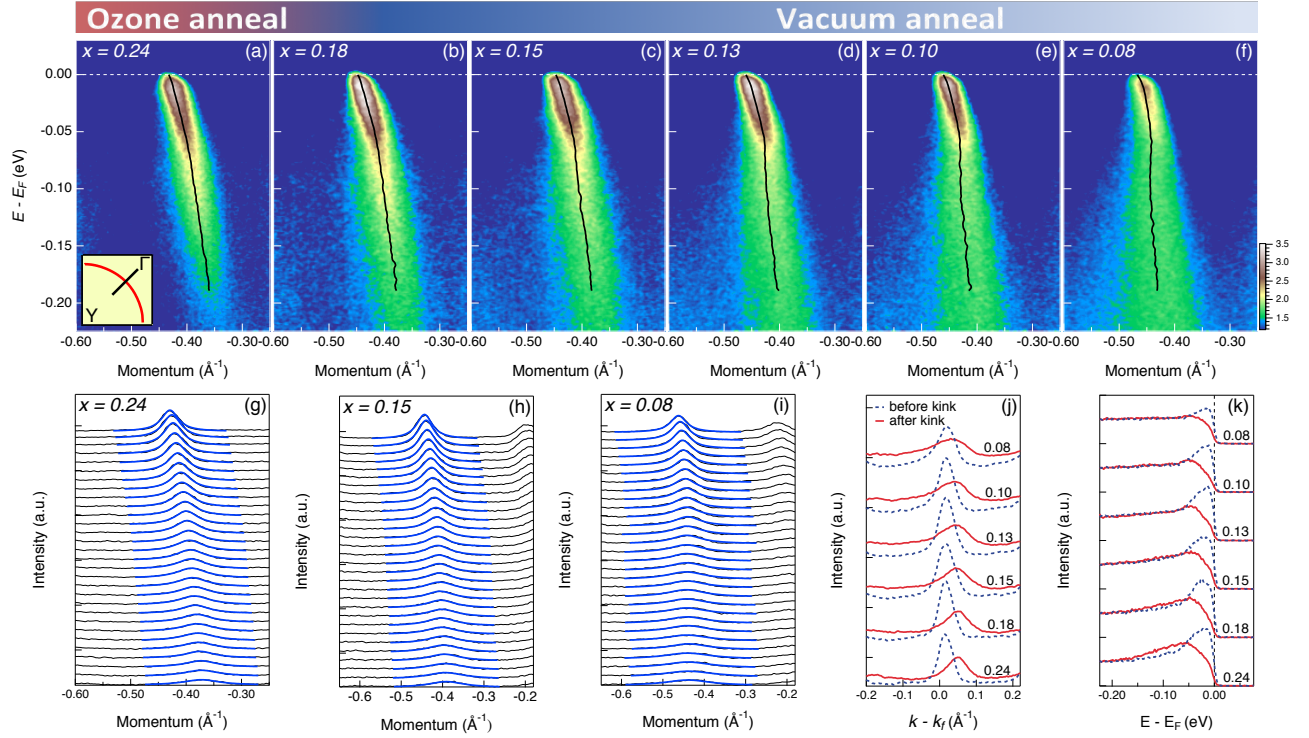


FIG. 1. (Color online) (a) ARPES spectrum of ozone-annealed Bi2212 along the nodal direction (as illustrated in the inset), the doping level is estimated as 0.24. Ozone anneal means sample is annealed at ozone atmosphere. (b) - (f) Spectra acquired from the same sample surface after annealed in vacuum with different temperatures (called vacuum anneal) step by step, and the corresponding doping levels are estimated as 0.18, 0.15, 0.13, 0.10, and 0.08, respectively. These spectra are plotted in the same color scale as illustrated in the lower right side of (f). (g) - (i) The black lines are the raw MDC plots for the doping levels at 0.24, 0.15 and 0.08, respectively, and the blue lines are their Lorentzian fittings. (j) The MDCs before the kink (blue dash lines,  $E - E_F = -27 \text{ meV}$ ) and after the kink (red solid lines,  $E - E_F = -105 \text{ meV}$ ) at different doping levels. (k) The EDCs before the kink (blue dash lines,  $k - k_f = 0.017 \text{ \AA}^{-1}$ ) and after kink (red solid line,  $k - k_f = 0.043 \text{ \AA}^{-1}$ ) at different doping levels.

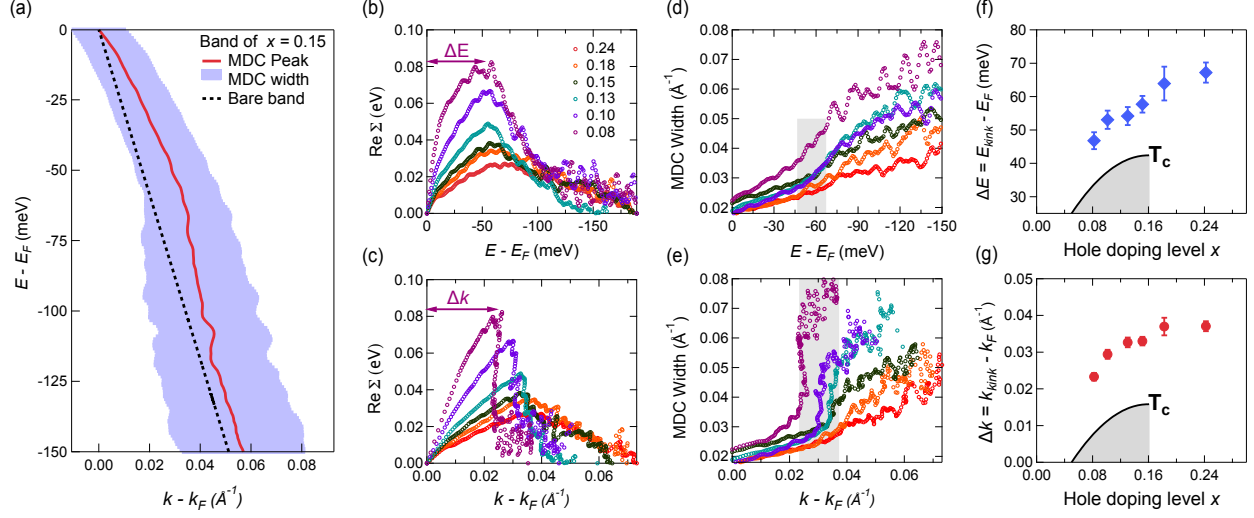


FIG. 2. (Color online) (a) MDC-derived band dispersion at the doping level equals 0.15: the red line is the peak of MDCs extracted by Lorentzian fitting, and the violet shadow represents the width of MDCs. The dashed line is a typical bare band as described in the text. (b)-(c) The real parts of SE at each doping level are plotted against with the binding energy in (b) and with the peaks' position of the corresponding MDCs. The double-headed arrows indicate how we extract the energy positions  $\Delta E(x)$  and momentum position  $\Delta k(x)$  respectively. (d)-(e) Same as (b) and (c) but for the MDCs width at each doping level. The shadow regions in (d) and (e) highlight the dramatic changes of the MDCs width. (f) - (g) The determined  $\Delta k(x)$  and  $\Delta E(x)$  for each doping level. The black lines along with the shadow domes represents the UD side of the superconducting dome.

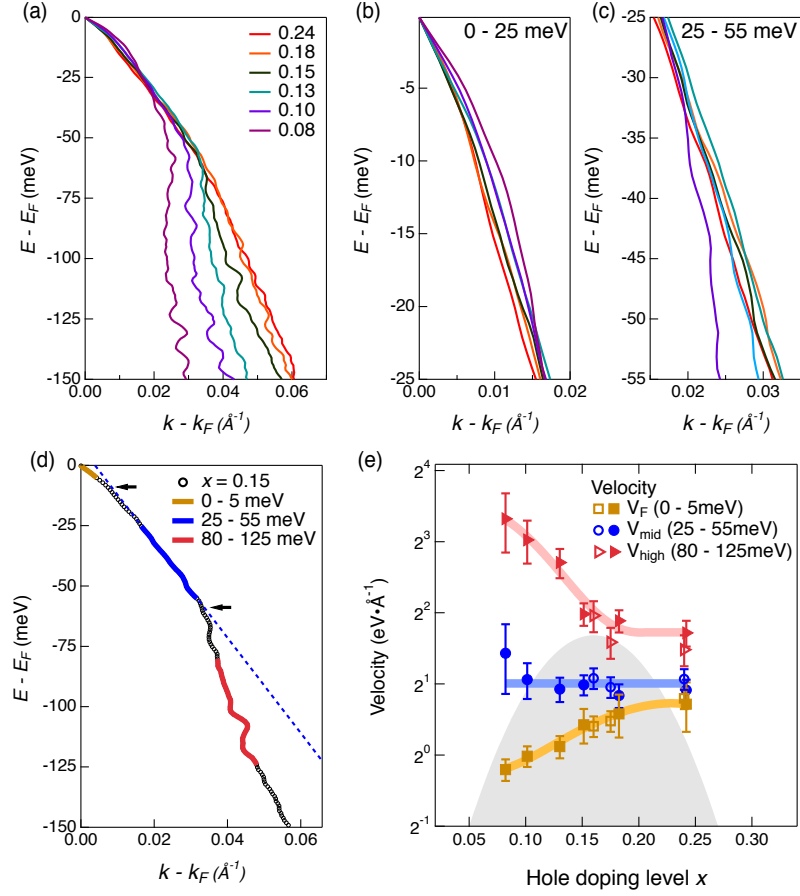


FIG. 3. (Color online) (a) Summary of MDC-derived nodal dispersions at different doping levels. The momentum is scaled to their corresponding  $k_F$ . (b) The MDC-derived nodal dispersion segments at the binding energy range with 0-25 meV. (c) The MDC-derived nodal dispersion segments at 25-55 meV. (d) Three segments separated by two kinks which are marked with the black arrows: the brown one marks the segment at the 0-5 meV, the blue one marks the segment at the 25-55 meV, the red one marks the segment at the 80-125 meV. The black circles are the nodal dispersion at the doping level equals 0.15. (e) The doping evolution of velocities of three segments plotted with the corresponding colors marked in (d). The empty markers with corresponding shapes represent an independent dataset acquired from another sample. The colored shadow lines are guides for eyes. The gray shadow dome represents the superconducting phase region.

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