

Temperature-Dependent Luttinger Surfaces

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The Luttinger surface of an organic metal (TTF-TCNQ), possessing charge order and spin-charge separated band dispersions, is investigated using temperature-dependent angle-resolved photoemission spectroscopy. The Luttinger surface topology, obtained from momentum distribution curves, changes from quasi-2D (dimensional) to quasi-1D with temperature. The high temperature quasi-2D surface exhibits $4k_F$ charge-density-wave (CDW) superstructure in the TCNQ derived holon band, in the absence of $2k_F$ order. Decreasing temperature results in quasi-1D nested $2k_F$ CDW order in the TCNQ spinon band and in the TTF surface. The results establish the link in momentum space between charge order and spin-charge separation in a Luttinger liquid.

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The Fermi surface defining a crystalline metal [1] is a set of points in momentum (k) space at which a dispersing quasiparticle band crosses the Fermi-level (E_F) in energy distribution curves (EDCs), with corresponding peaks in momentum distribution curves (MDCs) [2]. This holds for a correlated Fermi liquid. In contrast, a Luttinger liquid is a quasi-one-dimensional correlated metal with two important characteristics: (i) absence of quasiparticles and (ii) spin and charge separated excitations [3–8]. Thus, Luttinger liquids do not show quasiparticle band crossings at E_F in EDCs as measured by angle-resolved photoemission spectroscopy (ARPES) [3]. This is also true for many quasi-2D high- T_c cuprates and related materials [4–8]. For the cuprates, this has been explicitly shown to be due to electron fractionalization or spin-charge separation in a quasi-1D Luttinger liquid [5,6]. The spin and charge separated band dispersions of a Luttinger liquid are called spinon and holon bands, respectively [3,5]. Although they do not show quasiparticle band crossings, recent theory [9] has predicted that Luttinger liquids should exhibit a so-called “Luttinger surface”—the equivalent of a Fermi surface for a non-Fermi liquid, arising from spinon and holon bands.

Tetrathiafulvalene-tetracyanoquinodimethane (TTF-TCNQ), is an organic metal consisting of TTF (donor) and TCNQ (acceptor) columns stacked along the b axis, and has been extensively studied for its structural, transport, and thermodynamic properties [10–16]. It is a classic example of a quasi-1D correlated metal with a charge-density-wave (CDW) transition, exhibiting $2k_F$ and $4k_F$ CDW order along the b^* (or k_y) direction in k space [10–17]. These two aspects, namely, (i) the strong electron-electron correlations and (ii) the electron-phonon coupling induced CDW transition setting in at 54 K, are believed to occur in different parts of the electronic structure: the TTF electronic structure displays the $2k_F$ CDW order and the TCNQ part exhibits the $4k_F$ Coulomb correlation driven

order [10,11,16,18,19]. Early theory showed that the $2k_F$ and $4k_F$ CDW transition can occur in TTF-TCNQ, in the form of a correlated CDW state with lattice distortion, but required long-range Coulomb interactions [13]. More recently, TTF-TCNQ has also served as a very good example of a correlated metal with non-Fermi-liquid character, exhibiting spin-charge separated band dispersions [20,21]. The spin-charge separated band dispersions are consistent with 1D Hubbard model calculations in the absence of a temperature-dependent CDW transition [20,21]. In this work, we investigate the relation between spin-charge separated band dispersions and the temperature evolution of $2k_F$ and $4k_F$ CDW order involving specific electronic states constituting the TTF or TCNQ Luttinger surfaces. The results show that the $4k_F$ CDW superstructure is a special property of the holon band derived Luttinger surface.

Angle-resolved photoemission spectra were obtained from single crystals of TTF-TCNQ using a Scienta SES 2002 analyzer and a Gammadata-discharge lamp (He II α , $h\nu = 40.8$ eV). The energy and momentum resolutions for the ARPES measurements were 20 meV and 0.01 \AA^{-1} , respectively. A thin film filter [22] was instrumental in obtaining the spectra without degradation as a function of time and exposure to UV photons over several hours. The samples were cooled using a flowing liquid He cryostat and the temperature was controlled within ± 2 K using a temperature controller. Fresh surfaces were obtained by cleaving the samples *in situ* and the temperature dependence was confirmed to be reproducible.

In Figs. 1(a)–1(c) we plot two dimensional intensity maps as a function of momentum in the first Brillouin zone of TTF-TCNQ, for $T = 200, 100,$ and 30 K, obtained from ARPES spectra. The intensity maps are obtained for an energy window of 50 meV (E_F to 50 meV binding energy) and the high intensity regions correspond to Luttinger surfaces in the first Brillouin zone. At 200 K,

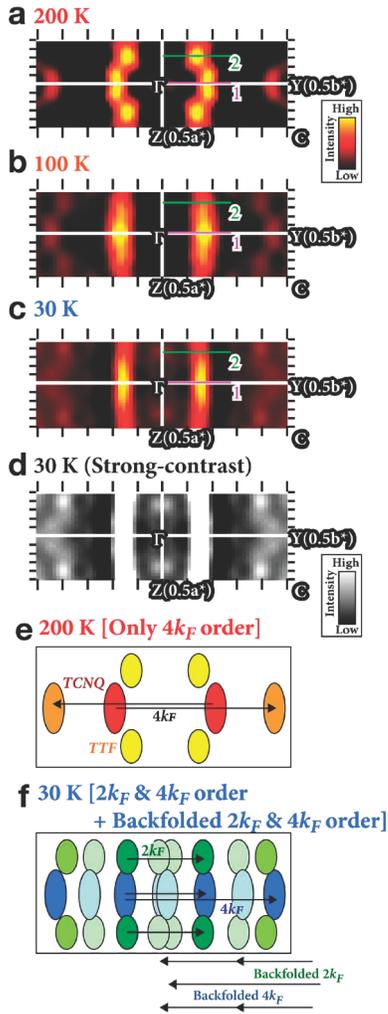


FIG. 1 (color). Temperature-dependent Luttinger surface maps: (a),(b),(c) show the Luttinger surfaces of TTF-TCNQ as a function of temperature (200, 100, and 30 K), obtained from angle-resolved photoemission spectra. (a) shows the TTF (along slice 2, centered at $k_y \sim \pm 0.13b^*$) and TCNQ (along slice 1, centered at $k_y \sim \pm 0.17b^*$) main Luttinger surfaces separated in momentum space [see also schematic (e)]. The TCNQ surface shows a $4k_F$ CDW superstructure at about $\pm 0.43b^*$ along slice 1 (Γ -Y) in k space. The $T = 100$ K map (b) shows features due to the $2k_F$ CDW order in addition to the $4k_F$ CDW vectors which evolve with reducing temperature. (c) shows that the low temperature ($T = 30$ K) main TTF and TCNQ Luttinger surfaces are nearly one dimensional, in good agreement with a nesting picture of the $2k_F$ CDW vector. The main Luttinger surfaces occur at $k_y \sim \pm 0.15b^*$ and the $4k_F$ superstructures are at $\sim \pm 0.45b^*$ along slice 1 (Γ -Y) in k space [dark blue and green regions shown schematically in (f)]. (d) is a gray scale map of (c), showing the weaker backfolded structures (at $k_y \sim 0b^*$ and $\pm 0.3b^*$) more clearly. Along slice 2, the superstructures reflect deviations from perfect nesting due to small differences in peak positions and widths of the main Luttinger surfaces. (e) schematically shows the presence of the $4k_F$ CDW order (connecting red and brown regions), in the absence of $2k_F$ CDW order. Light blue and green regions in schematic (f) show the relation of the backfolded features with the $2k_F$ and $4k_F$ CDW order.

the main Luttinger surfaces consisting of the TCNQ surface (along slice 1, centered at $k_y \sim \pm 0.17b^*$) and the TTF surface (along slice 2, centered at $k_y \sim \pm 0.13b^*$), are well separated in k space and are quasi-2D. These are shown schematically in Fig. 1(e) as red (TCNQ) and yellow (TTF) regions. Additional replica features only for the TCNQ branch are seen in Fig. 1(a) at momenta translated by a value of $\sim 0.6b^*$, along slice 1 [brown regions in schematic Fig. 1(e)]. The observation of replica features translated by $\sim 0.6b^*$ correspond to the observation of $4k_F$ order, consistent with superstructure spots in structural studies [12,14,15]. This surface arises from the holon band crossing at $4k_F$, which is absent in band structure calculations [23], but is obtained in the 1D Hubbard model [20,21]. While the high energy band dispersions have been reported for the TCNQ holon band [20], a Luttinger surface obtained from low energy spectral weight (within 50 meV of E_F) has not been possible earlier due to the very low intensity near E_F . The holon band derived Luttinger surface is consistent with the 1D Hubbard model at low energies. Its presence also indicates that the main TCNQ band along slice 1 is the spinon band and not simply the uncorrelated TCNQ band expected from band structure calculations. Band structure calculations show that the TCNQ surface occurs along slice 1 and the TTF surface occurs along slice 2 in k space [23]. We have measured and confirmed the high energy dispersions (not shown) to be in excellent agreement with earlier work [19,20]. At $T = 200$ K, since the separation between the main TTF and TCNQ bands are not equal to $2k_F$, the $4k_F$ order exists in the absence of $2k_F$ order. The $2k_F$ CDW order is expected to show up only below 150 K, as is known from earlier studies [12,14,15].

At $T = 100$ K, the separation along the b^* direction seen in the TTF and TCNQ branches is reduced and they merge together [Fig. 1(b)], with the observation of additional weak features in the map. These changes occur above the CDW transition which sets in below 54 K and correspond to the growth of the $2k_F$ CDW order. While these weak features derived from the low energy electronic structure were not observed in earlier ARPES studies [19,20], they are consistent with a recent analysis of x-ray diffuse scattering above the $2k_F$ CDW transition, due to quasi-1D structural fluctuations [24]. At the lowest temperature of 30 K [Fig. 1(c)], below the CDW transition at 54 K, followed by additional order at 38 K, the TTF and TCNQ parts of the main Luttinger surfaces satisfy the nesting condition for the $2k_F$ CDW order and are nearly one dimensional.

MDCs obtained at $T = 30$ K with high signal to noise ratio, using the same energy window of 50 meV as for the maps of Fig. 1, are shown along Γ -Y [Fig. 2(a), slice 1] and along Z-C [Fig. 2(b), slice 2]. The MDC curves show sharp peaks at equidistant points in k space from the Γ [Fig. 2(a)] and Z points [Fig. 2(b)] at a value of $\sim (\pm 0.15 \pm 0.01)b^*$.

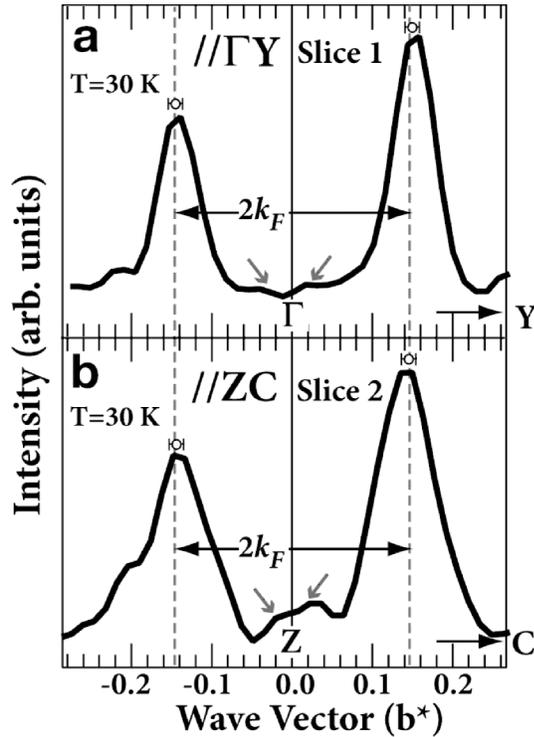


FIG. 2. Momentum distribution curves of TTF-TCNQ: (a),(b) show the momentum distribution curves (MDCs) along slice 1 (TCNQ derived band) and slice 2 (TTF-derived band) at $T = 30$ K obtained using ARPES spectra. At $T = 30$ K, the separation in their positions $(0.3 \pm 0.01)b^*$ matches the CDW vector ($2k_F = 0.295b^*$) known from structural studies. The peak widths imply a CDW coherence length of ~ 120 Å for the TCNQ and ~ 60 Å for the TTF-derived surfaces. Small gray arrow marks the weak superstructure also seen in the maps [Fig. 1(d)].

The peak-to-peak separation in k space ($\sim 0.30b^*$) matches the CDW vector ($2k_F = 0.295b^*$) in TTF-TCNQ, known from structural studies [12,14,15]. Similar peaks in MDCs have been observed in the quasi-1D conductors $K_{0.3}MoO_3$ [25] and $(TaSe_4)_2I$ [26], which undergo the Peierls' transition but do not show evidence for spin-charge separation. The MDC peak widths at $T = 30$ K for the TCNQ spinon band derived surface is $\sim 0.05b^* \text{ \AA}^{-1}$ [Fig. 2(a)]. This peak width implies a CDW coherence length of ~ 120 Å. The MDC peak width [$\sim 0.1b^* \text{ \AA}^{-1}$, Fig. 2(b)] for the TTF-derived surface implies a shorter CDW coherence length of ~ 60 Å. These values are consistent with recent x-ray diffuse scattering studies [24] and suggest that the TCNQ derived CDW order is more robust and induces the CDW order in the TTF stacks. The results indicate that the $2k_F$ and $4k_F$ order in the TCNQ surface is derived from the correlation induced spinon and holon bands, respectively, and the TTF surface has only the $2k_F$ CDW order. The $2k_F$ CDW order involves electron-phonon coupling, as is known from the Kohn anomaly observed by neutron scattering [14]. This clarifies the specific role of TTF and TCNQ derived electronic states as a function of

momentum and temperature compared with earlier work [11,12,14,15,27].

In addition to the main TTF and TCNQ Luttinger surfaces, additional weak features are seen at $T = 30$ K in the MDCs of Figs. 2(a) and 2(b), marked with small arrows. These features are better seen in Fig. 1(d), which is a gray scale image of Fig. 1(c), plotted in order to see the weak features with stronger contrast. The $T = 30$ K gray scale image shows many more features compared to the 200 K plot [Fig. 1(a)]. We checked for the separation between the features along b^* and find that the weak features occur exactly at multiples of $2k_F$ and $4k_F$ vectors obtained by a backfolding from the second Brillouin zone. The deviations from quasi-1D behavior in the weak features is attributed to the difference in peak widths and positions along b^* in the main TTF and TCNQ Luttinger surfaces, which get multiplied for the weak features. A very weak feature is also seen in Fig. 1(a) near the Γ point due to a backfolding of the $4k_F$ CDW vector at $T = 200$ K. A schematic of the high- and low- T main Luttinger surfaces and the backfolded Luttinger surfaces is drawn in Figs. 1(e) and 1(f) to explain all the features observed in the maps. Figure 1(e) shows schematically the main Luttinger surfaces of TTF (yellow regions), and TCNQ (red regions) along with the $4k_F$ CDW superstructure (brown regions) at $T = 200$ K. At $T = 30$ K, Fig. 1(f) schematically shows the main (dark blue and green regions) and backfolded (light blue and green regions) superstructures due to the $2k_F$ and $4k_F$ CDW order.

The observation of temperature-dependent Luttinger surfaces indicates that angle-resolved photoemission spectroscopy provides important insights about CDW order in a Luttinger liquid, and its spectral function retains Fermi-liquid-like properties in terms of peaks in MDCs, although the quasiparticles in EDCs are suppressed [3–9,19]. The evolution from a Mott-Insulator (MI) to Luttinger liquid and then on to a Fermi liquid with a concomitant transformation of the Fermi surface topology was predicted theoretically for the Bechgaard salts, in a model of correlated chains with interchain hybridization [28]. Optical measurements indeed reveal evidence for spin-charge separation in the Bechgaard salts and a dimensionally driven insulator-metal transition [29]. Luttinger surface studies are eagerly awaited for this family of quasi-1D correlated metals which also exhibit density-wave order and superconductivity. If we identify the MI phase with the $2k_F$ CDW ordered phase, the same theoretical phase diagram [28] holds for the topology change observed in TTF-TCNQ [Figs. 1(a)–1(c)]. While recent work has [30,31] identified the power law characteristic of Luttinger liquids in angle-integrated photoemission spectra of carbon nanotubes [30,31], and also a doping dependent Luttinger to Fermi-liquid crossover [31], ARPES of the Fermi or Luttinger surfaces in the carbon nanotube systems are intrinsically difficult.

In conclusion, the Luttinger surface topology, obtained from MDCs, changes from quasi-two-dimensional to quasi-one-dimensional with temperature. The high temperature quasi-2D surface exhibits $4k_F$ CDW superstructure in the TCNQ derived holon band, in the absence of $2k_F$ order. On decreasing temperature, a quasi-1D nested $2k_F$ CDW order in the TCNQ spinon band and in the TTF surface is obtained. The results show the link in momentum space between charge order and spin-charge separated band dispersions in a Luttinger liquid. Thus, the $4k_F$ CDW superstructure is a special property of the holon band derived Luttinger surface. The present study indicates T -dependent Luttinger surface topology is very important for density-wave phenomena in low-dimensional correlated systems.

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- [1] N.W. Ashcroft and N.D. Mermin, *Solid State Physics* (Saunders, Florida, 1976), p. 142.
- [2] T. Valla, A.W. Fedorov, P.D. Johnson, B.O. Wells, S.L. Hulbert, Q. Li, G.D. Gu, and N. Koshizuka, *Science* **285**, 2110 (1999).
- [3] J.W. Allen, *Solid State Commun.* **123**, 469 (2002).
- [4] Y. Ren and P.W. Anderson, *Phys. Rev. B* **48**, 16662 (1993).
- [5] C. Kim, A.Y. Matsuura, Z.-X. Shen, N. Motoyama, H. Eisaki, S. Uchida, T. Tohyama, and S. Maekawa, *Phys. Rev. Lett.* **77**, 4054 (1996); M.Z. Hasan, P.A. Montano, E.D. Isaacs, Z.-X. Shen, H. Eisaki, S.K. Sinha, Z. Islam, N. Motoyama, and S. Uchida, *Phys. Rev. Lett.* **88**, 177403 (2002).
- [6] D. Orgad, S.A. Kivelson, E.W. Carlson, V.J. Emery, X.J. Zhou, and Z.-X. Shen, *Phys. Rev. Lett.* **86**, 4362 (2001).
- [7] G.H. Gweon, J.W. Allen, and J.D. Denlinger, *Phys. Rev. B* **68**, 195117 (2003).
- [8] S.A. Kivelson, I.P. Bindloss, E. Fradkin, V. Oganessian, J.M. Tranquada, A. Kapitulnik, and C. Howald, *Rev. Mod. Phys.* **75**, 1201 (2003).
- [9] I. Dzyaloshinskii, *Phys. Rev. B* **68**, 085113 (2003).
- [10] A.J. Heeger, in *Highly Conducting One-Dimensional Solids*, edited by J. Devreese (Plenum, New York, 1979) p. 69.
- [11] Y. Tomkiewicz, A.R. Taranko, and J.B. Torrance, *Phys. Rev. Lett.* **36**, 751 (1976).
- [12] J.P. Pouget, S.K. Khanna, F. Denoyer, R. Comes, A.F. Garito, and A.J. Heeger, *Phys. Rev. Lett.* **37**, 437 (1976).
- [13] V.J. Emery, *Phys. Rev. Lett.* **37**, 107 (1976); P. Bak and V.J. Emery, *Phys. Rev. Lett.* **36**, 978 (1976); V.J. Emery, A. Luther, and I. Peschel, *Phys. Rev. B* **13**, 1272 (1976).
- [14] G. Shirane, S.M. Shapiro, R. Comes, A.F. Garito, and A.J. Heeger, *Phys. Rev. B* **14**, 2325 (1976).
- [15] S. Kagoshima, H. Nagasawa, and T. Sambongi, *One-Dimensional Conductors* (Springer-Verlag, Berlin, 1987).
- [16] G. Gruner, *Density Waves in Solids* (Addison-Wesley, Reading, 1994).
- [17] S. Brazovskii, cond-mat-0502264 [Synth. Met. (to be published)].
- [18] D. Jerome, *Chem. Rev.* **104**, 5565 (2004).
- [19] F. Zwick, D. Jerome, G. Margaritondo, M. Onellion, J. Voit, and M. Grioni, *Phys. Rev. Lett.* **81**, 2974 (1998).
- [20] R. Claessen, M. Sing, U. Schwingenschlögl, P. Blaha, M. Dressel, and C. Jacobsen, *Phys. Rev. Lett.* **88**, 096402 (2002).
- [21] M.P. Carmelo, K. Penc, L.M. Martelo, P.D. Sacramento, J.M.B. Lopes dos Santos, R. Claessen, M. Sing, and U. Schwingenschlögl, *Europhys. Lett.* **67**, 233 (2004).
- [22] T. Yokoya, S. Tsuda, T. Kiss, S. Shin, T. Mochiku, and K. Hirata, *Jpn. J. Appl. Phys.* **43**, 3618 (2004).
- [23] S. Ishibashi and M. Kohyama, *Phys. Rev. B* **62**, 7839 (2000).
- [24] J.P. Pouget, *Z. Kristallogr.* **219**, 711 (2004).
- [25] A.V. Fedorov, S.A. Brazovskii, V.N. Muthukumar, P.D. Johnson, J. Xue, L.-C. Duda, K.E. Smith, W.H. McCarroll, M. Greenblatt, and S.L. Hulbert, *J. Phys. Condens. Matter* **12**, L191 (2000).
- [26] L. Perfetti, H. Berger, A. Reggiani, L. Degiorgi, H. Höchst, J. Voit, G. Margaritondo, and M. Grioni, *Phys. Rev. Lett.* **87**, 216404 (2001).
- [27] T. Takahashi, D. Jerome, F. Masin, J. Fabre, and L. Giral, *J. Phys. C* **17**, 3777 (1984).
- [28] S. Biermann, A. Georges, A. Lichtenstein, and T. Giamarchi, *Phys. Rev. Lett.* **87**, 276405 (2001).
- [29] V. Vescoli, L. Degiorgi, W. Henderson, G. Gruner, K.P. Starkey, and L.K. Montgomery, *Science* **281**, 1181 (1998).
- [30] H. Ishii, H. Kataura, H. Shiozawa, H. Yoshioka, H. Otsubo, Y. Takayama, T. Miyahara, S. Suzuki, Y. Achiba, M. Nakatake, T. Narimura, M. Higashiguchi, K. Shimada, H. Namatame, and M. Taniguchi, *Nature (London)* **426**, 540 (2003).
- [31] H. Rauf, T. Pichler, M. Knupfer, J. Fink, and H. Kataura, *Phys. Rev. Lett.* **93**, 096805 (2004).