Comment on "Fermi Surface, Surface States, and Surface Reconstruction in Sr₂RuO₄"

In a recent Letter, Damascelli et al. [1] reported angleresolved photoemission spectra on Sr_2RuO_4 with the aim of elucidating the electronic structure in the vicinity of the Fermi energy. They interpret their data in terms of a surface state just below E_F near the M point, shadow bands induced by the $\sqrt{2} \times \sqrt{2}$ surface reconstruction, and Ru $4d t_{2g}$ bands that are consistent with de Haas-van Alphen (dHvA) measurements [2]. Although these results resolve the discrepancy which existed between earlier photoemission spectra and the Fermi surface topology derived from dHvA data, we point out that the bulk band structure of Sr₂RuO₄ [3] provides no evidence of a band gap supporting the existence of a surface state. Thus the interpretation given in [1] of the prominent emission feature near M in terms of a surface state localized in k space cannot be correct. A clarification of the origin of this feature, however, is crucial since it sheds light on the usefulness of angle-resolved photoemission as a probe of Fermi surface electronic properties and since the controversial $t_{2,a}$ band in Sr₂RuO₄ shares important characteristics with the key electronic band in the high- T_c superconductors. Also, a clear picture of the electronic properties of Sr₂RuO₄ is desirable in view of the fact that this system is the only layered perovskite without copper that exhibits superconductivity [4] and appears to show unconventional spin-triplet pairing with a *p*-wave order parameter [5].

An interpretation of the observed photoemission spectra that is consistent with all available experimental and theoretical results can be achieved by associating the feature near M with emission from the deformed xy band within the reconstructed first layer of Sr₂RuO₄. According to recent surface electronic structure calculations, scanning tunneling microscopy and low energy electron diffraction data [6], the following picture emerges for the properties of the first layer: The reconstruction is driven by the freezing of a zone boundary soft phonon mode giving a slight rotation of the RuO₆ octahedra around the surface normal. The rotation reduces the effective d-d hopping within the first Ru plane and causes band narrowing. In the nonmagnetic case (which seems to be observed in the photoemission work) the band narrowing yields a shift of the xy Van Hove singularity (VHs) below E_F , giving rise to strong emission near M.

The photoemission spectra in this picture consist of a superposition of two sets of t_{2g} bands originating in the surface layer and the deeper layers. Both sets are energetically very similar and difficult to resolve, with the exception of the xy VHs near M: in the bulk it is above E_F while at the surface it is below. Accordingly, the γ sheet of the Fermi surface is electronlike in the bulk, but holelike at the surface. Whereas the bulk xy band crosses the Fermi level

along ΓM , the reconstruction-induced *xy* band remains below E_F near *M* and crosses along *MX*. Both bands cross the Fermi level along ΓX . In addition, the reconstruction generates weak shadow bands, as observed in [1].

The weak momentum dispersion of the xy band near M can be understood in terms of Coulomb correlations between Ru d electrons. As shown recently using perturbation theory and quantum Monte Carlo calculations for a multiband Hamiltonian [7], an on-site Coulomb energy $U \approx 1$ eV leads to a quasiparticle self-energy near E_F of the same order as the binding energy, giving rise to considerable flattening of the momentum dispersion and strong effective mass enhancement. Correlations also cause a slight charge transfer from the narrow xz, yz bands to the wider xy band, strong quasiparticle damping, and an overall narrowing of the xz, yz bands by about a factor of 2, in agreement with photoemission data. None of these aspects can be understood within the single-particle picture and they demonstrate the importance of considering correlation effects when interpreting photoemission spectra.

The sensitivity of the xy band to surface degradation is at first sight surprising since adsorption of atoms or molecules most likely involves primarily xz, yz states rather than planar xy states. Nevertheless, adsorption requires reorganization of all Sr 6s, O 2p, and Ru 4d electrons in the first layer. It is plausible that this affects also the VHs near M. Quantitative chemisorption calculations might help to clarify this point.

In conclusion, the Sr₂RuO₄ photoemission spectra of Damascelli *et al.* [1] are interpreted in terms of a superposition of bulk t_{2g} states consistent with dHvA measurements and deformed t_{2g} states existing in the reconstructed first layer. The main deformation arises near *M* where the VHs of the *xy* band is below E_F as a result of a slight narrowing of the t_{2g} bands, while the bulk VHs is unoccupied.

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Damascelli et al. Reply: In our Letter [1] we showed that, in order to gain direct insight on the bulk electronic structure of Sr₂RuO₄ by angle-resolved photoemission spectroscopy (ARPES), it is necessary to separate the bulk from the surface contributions to the total photoemission intensity. In particular, only when the surface signal has been suppressed by means of a controlled degradation of the sample surface are the ARPES results representative of the bulk and is the corresponding Fermi surface in agreement with the results of band structure calculations [2] and de Haas-van Alphen experiments [3]. Most of the surface signal could be described in terms of the rigid folding of the bulk electronic structure with respect to the M-Mdirection [1], consequent of the (previously overlooked) $\sqrt{2} \times \sqrt{2}$ surface reconstruction [1,4]. However, this simplistic scheme failed to account for the intense feature observed at the M point, which led us to consider a different origin for the latter surface band (including, e.g., surface ferromagnetism [1,4,5]).

In his Comment [6] Liebsch points out that, once the distortion of the surface electronic structure due to the reduced symmetry is taken into account (beyond the naive rigid band folding), the ARPES results from Sr₂RuO₄ could be interpreted purely as the superposition of bulk and reconstructed-surface contributions, without assuming the presence of any additional surface states. We here emphasize that Liebsch's interpretation is not only plausible, but indeed appears to be correct, as also evidenced by an extensive experimental and theoretical investigation of the surface electronic structure performed by our group [7]. By comparing a more complete body of experimental data with band structure calculations for the reconstructed surface (with and without ferromagnetism), we could ascribe all the surface features detected by ARPES to a nonmagnetic reconstructed surface (as independently suggested by Liebsch [6]) and rule against the presence of additional surface states and/or surface ferromagnetism [7]. For example, along M-X, in addition to the surface bands at the M point already discussed in Refs. [1,6], a surface α pocket and a surface band located $\sim 3 \text{ meV}$ above E_F at X were clearly resolved in these more recent ARPES experiments [7]. As summarized in Fig. 1a [7], all these features are described in great detail by band structure calculations for the nonmagnetic reconstructed surface, while no match is found when both surface reconstruction and ferromagnetism are considered (Fig. 1b).

Liebsch further remarks [6] that, in order to properly interpret the ARPES results from Sr_2RuO_4 , the effect of electronic correlations must be taken into account [8]. This would naturally explain the experimentally observed quasiparticle renormalization although, contrary to what was previously suggested [8], it is not essential to resolve the Fermi surface topology controversy, which can be fully understood within an independent particle picture [7].



FIG. 1. (a) Band structure calculations for bulk and nonmagnetic reconstructed surface along M-X (thick and thin lines) and corresponding bulk and surface bands as detected by ARPES (solid and open circles). (b) Calculated electronic structure for the bulk (thick lines) and a ferromagnetic reconstructed surface (thin and dashed lines for majority and minority spin populations, respectively). After Shen *et al.* [7].

However, given the relatively small value of $U \approx 1 \text{ eV}$ needed in the multiband quasiparticle calculations [8] in order to reproduce the experimentally observed renormalization, we do not feel that Sr₂RuO₄ represents a clear-cut case for electron correlations as opposed to magnetic correlations [9].

Last, we point out that all the surface bands exhibit, in the ARPES experiments, comparable sensitivity to surface degradation [7]. This together with scanning tunneling microscopy results [10] suggest that cleaving the samples at elevated temperatures introduces substantial oxygen deficiencies at the surface which, in turn, would reduce the surface contribution to the photoemission signal to an angle independent background. This scenario however requires further scrutiny.

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