Molecular orbital excitations in cuprates

Young-June Kim, J. P. Hill, and G. D. Gu
Department of Physics, Brookhaven National Laboratory, Upton, New York 11973

F. C. Chou
Center for Materials Science and Engineering, Massachusetts Institute of Technology, Cambridge, Massachusetts 02139

S. Wakimoto and R. J. Birgeneau
Department of Physics, University of Toronto, Toronto, Ontario M5S 1A7, Canada

Seiki Komiya and Yoichi Ando
Central Research Institute of Electric Power Industry, Komae, Tokyo 201-8511, Japan

N. Motoyama, K. M. Kojima, and S. Uchida
Graduate School of Frontier Sciences, University of Tokyo, Bunkyo, Tokyo 113-8656, Japan

D. Casa and T. Gog
CMC-CAT, Advanced Photon Source, Argonne National Laboratory, Argonne, Illinois 60439
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We report resonant inelastic x-ray scattering studies of electronic excitations in a wide variety of cuprate compounds. Specifically, we focus on the charge-transfer type excitation of an electron from a bonding molecular orbital to an antibonding molecular orbital in a copper oxygen plaquette. Both the excitation energy and the amount of dispersion are found to increase significantly as the copper oxygen bond-length is reduced. We also find that the estimated bond-length dependence of the hopping integral $t_{pd}$ is much stronger than that expected from tight-binding theory.

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I. INTRODUCTION

One of the most important characteristics of the electronic structure of the cuprates is the strong hybridization between the Cu 3d_{x^2-y^2} level and the O 2p level, where $p_\sigma$ denotes the $p_x$ or $p_y$ orbitals pointing towards the Cu ions. Because of this hybridization, the Cu-O bond has a strong covalent character and a large energy splitting exists between the bonding ($\sigma$) and antibonding ($\sigma^*$) molecular orbitals. In the ionic limit without hybridization, this energy splitting corresponds to the energy difference between the atomic $d_{x^2-y^2}$ and $p_\sigma$ orbitals, which is $\Delta_0 \sim 3.5$ eV. As the $p-d$ hybridization becomes larger, the energy splitting ($\Delta_{\sigma\sigma^*}$) between the two molecular orbitals increases, reflecting the increasingly covalent nature of the Cu-O bonding. Thus, $\Delta_{\sigma\sigma^*}$ is a direct measure of the Cu-O hybridization, and could serve as an independent route to determine the value of the hopping matrix element, $t_{pd}$, since it is believed that $\Delta_{\sigma\sigma^*}$ is directly related to $t_{pd}$. Although there has been no systematic theoretical study of $\Delta_{\sigma\sigma^*}$, the values obtained from first-principles calculations range widely from ~4 eV to ~9 eV. For example, Martin and Hase have carried out an $ab$ initio quantum chemistry calculation of a cluster of copper oxygen octahedra (CuO$_6$) in La$_{1.85}$Sr$_{0.15}$CuO$_4$, and obtained $\Delta_{\sigma\sigma^*} \sim 9$ eV, while in a recent density-functional calculation of a similar cluster, Husser and coworkers reported a value of $\Delta_{\sigma\sigma^*} \sim 5.8$ eV. Experimentally, determining this quantity has been very difficult, since in this energy range transitions involving the La 4f bands dominate the spectral features of optical spectroscopy.

In this paper, we report a systematic experimental study of $\Delta_{\sigma\sigma^*}$, the excitation energy from bonding to antibonding molecular orbitals, using the recently developed resonant inelastic x-ray scattering (RIXS) technique. RIXS is ideally suited for this study, since it provides element-specific and momentum-dependent information for electronic excitations. By tuning the incident photon energy to the Cu-K absorption edge, one can gain information concerning excitations localized around the Cu sites without suffering from problems due to the La bands. Furthermore, the momentum-resolving capability of RIXS provides additional information: the dispersion of such molecular orbital (MO) excitations. In this work, we find that $\Delta_{\sigma\sigma^*}$ exhibits a strong, systematic dependence on the Cu-O bond length ($d_{Cu-O}$), increasing as $d_{Cu-O}$ is decreased. In addition, for materials with a small $d_{Cu-O}$ and correspondingly large $\Delta_{\sigma\sigma^*}$, a relatively large dispersion of the MO excitation is observed. We discuss the implication of these observations for understanding the electronic structure of the cuprates.

II. EXPERIMENTS

The RIXS experiments were carried out at the Advanced Photon Source on the undulator beamline 9IDB.
TABLE I: The copper oxide samples are listed along with the Cu-O bond lengths taken from the references. The top eight materials possess perfect square copper oxygen plaquettes, while the bottom three have distorted square plaquettes. Also listed are the polarization and the energy of the incident photon in RIXS measurements.

| Label | Sample | Crystal | $d_{\text{Cu-O}}$ (Å) | Ref. ($d_{\text{Cu-O}}$) | Polarization | $E_r$ (eV) | Ref. (RIXS) |
|-------|--------|---------|------------------------|--------------------------|--------------|------------|------------|
| 2122  | Sr$_2$CuO$_2$Cl$_2$ | M | 1.9858 | 12 | $\epsilon \perp z$ | 9001 | This work |
| Nd    | Nd$_2$CuO$_4$ | | 1.9705 | 13 | $\epsilon \perp z$ | 8990 | 14 |
| Ca    | Ca$_2$CuO$_2$Cl$_2$ | | 1.9344 | 15 | $\epsilon \parallel z$ | 8996 | 16 |
| 2342  | Sr$_2$Cu$_3$O$_4$Cl$_2$ | M | 1.929 | 17 | $\epsilon \perp z$ | 8998 | This work |
| LCCO  | La$_{1.9}$Ca$_{1.1}$Cu$_2$O$_6$ | B | 1.913 | 18 | $\epsilon \parallel z$ | 8999 | This work |
| LCO   | La$_2$CuO$_4$ | T | 1.904 | 19 | $\epsilon \parallel z$ | 8997 | This work |
| LSCO5 | La$_{1.95}$Sr$_{0.05}$CuO$_4$ | C | 1.898 | 20 | $\epsilon \parallel z$ | 8997 | This work |
| LSCO17| La$_{1.83}$Sr$_{0.17}$CuO$_4$ | C | 1.885 | 21 | $\epsilon \parallel z$ | 8997 | 21 |
| Li    | Li$_2$CuO$_2$ | | 1.9577 | 22 | $\epsilon \parallel z$ | 8990 | 23 |
| CGO   | CuGeO$_3$ | | 1.9326 | 24 | $\epsilon \parallel z$ | 8996 | This work |
| 112   | Sr$_2$CuO$_4$ | U | 1.910/1.930/1.961 | $d$ | $\epsilon \parallel z$ | 8996 | This work |

*The crystals studied in this work were provided by various groups, which are denoted here as B:Brookhaven; C:CREPI; M:MIT; T:Toronto; U:Univ. of Tokyo.

Since the polarization direction was in the scattering plane in this experiment, it changed as momentum transfer was varied.

The RIXS data were taken at $T = 15$ K.

Sr$_2$CuO$_2$ has three different copper-oxygen bond lengths, which are represented as large error bars in Fig. 2.

Experimental details have been described elsewhere. Single crystal samples used in our measurements are listed in Table I along with several samples studied in earlier RIXS experiments. In Table I, $d_{\text{Cu-O}}$ and the experimental configuration is listed for each material. All measurements were performed at room temperature except for those on the LSCO17 sample. In our RIXS experiments, the scattering plane was vertical and the polarization of the incident x-ray, $\epsilon$, was perpendicular to the scattering plane. The polarization direction was kept fixed along the direction specified in Table I where the coordinate system reference is the $d_{\text{xy}}$ direction along the Cu-O bond direction.

Before discussing the experimental results in detail, it is useful to first review the second-order RIXS process to understand the nature of the observed excitation. In the ground state, the holes are located in the antibonding molecular orbital which is a combination of a Cu hole state ($d^9$) and an oxygen ligand hole state ($d^{10}L$), with more weight on the $d^9$ state. In the intermediate state of this resonance process, a Cu 1s electron is excited to the Cu 4p band, and the core hole potential alters the balance between the $d^9$ and the $d^{10}L$ states. Then the lowest energy state is predominantly $1s_{\text{Cu}}d^{10}4p$, which is lower than $1s_{\text{Cu}}d^{9}4p$. These states form the so-called well-screened and poorly-screened features, respectively, of the Cu K-edge x-ray absorption spectra (XAS). As discussed in detail by Hill and coworkers, these intermediate states can decay into an excited state in which the hole in the antibonding molecular orbital is filled with an electron, creating a hole in the bonding orbital, and an energy loss in the outgoing photon. The RIXS process thus creates a charge-transfer excitation from bonding to antibonding molecular orbitals.

In our measurements, we have carefully studied the incident energy dependence in order to determine the resonance energy, for which the MO excitation has the maximum intensity. In most cases, the resonance energy, which is listed in Table I, corresponds to the higher energy peak (poorly-screened feature) in the XAS. On the other hand, the intensity of lower energy excitations near the charge transfer gap ($\sim 2$ eV) is strongly enhanced when the incident energy corresponds to the well-screened intermediate state, as reported in earlier studies. The resonance energy depends on the direction of the polarization vector. Detailed results of the incident energy dependence study will be published elsewhere. However, one should note that the energy-loss associated with the MO excitation does not depend on the incident polarization of the photon, although different 4p states (e.g., 4p$_x$ or 4p$_z$) are involved in the intermediate state, as the polarization is varied with respect to the $xy$-plane.

III. RESULTS

In Fig. I, representative RIXS scans are plotted. These are energy-loss scans taken at a fixed momentum transfer with the incident energy of the x-ray photon fixed at the values listed in Table I. The momentum transfer for all these scans has been fixed at $q = (\pi, 0)$, which is the minimum energy position. The most striking feature in Fig. I is the large shift of the excitation energy from $\sim 6$ eV for...
FIG. 1: RIXS spectra taken with the incident energy as specified in Table I for a fixed reduced wave vector of \((\pi, 0)\). Each spectrum is offset vertically for clarity, and solid lines are fits to a Lorentzian lineshape as described in the text. The dashed line is a representative scan through the elastic line, which shows instrumental energy resolution.

2122 to \(\sim 8\) eV for LSCO17. To analyze this shift quantitatively, we have fitted the observed excitation spectra to a simple Lorentzian lineshape and extracted the peak positions, which are plotted in Fig. 2(a) as a function of Cu-O bond length, \(d_{\text{Cu-O}}\).

A few comments are in order regarding the data analysis. First, as is evident from the instrumental resolution plotted as a dashed line in Fig. 1, the observed excitations are not resolution-limited, hence justifying our simple fitting procedure, which does not convolve the data with the instrumental resolution. Second, in several cases, we observe more than one type of excitation in these scans. For example, the SrCuO$_2$ data clearly shows a second feature around \(\sim 8.2\) eV, in addition to the main peak around \(\sim 6\) eV. In the 2342 case, there also seems to be two additional features, one at higher energy (\(\sim 8\) eV) and the other at lower energy (\(\sim 4\) eV). However, it is difficult to identify these weak features (if present) in the data for the other samples and we have chosen to fit all the scans with a single Lorentzian peak with a broad width. The peak positions extracted from our analysis are, therefore, those of the dominant features. In addition, the peak width extracted from the fits might in some cases arise from a distribution of several peaks over a wide energy range, rather than from a finite inverse lifetime of a single excitation. Finally, as discussed below, we observe dispersion of the MO excitation with momentum transfer. This can be as large as \(\sim 0.5\) eV in some of the compounds studied, as shown in Fig. 3. Thus, the peak position of the MO excitation depends not only on the sample, but also on \(q\). The scans shown in Fig. 1 are taken at the minimum energy position, \(q = (\pi, 0)\), and the peak positions plotted in Fig. 2(a) are the values measured at this position.

As shown in Fig. 2(a), the energy of the MO excitation exhibits a strong dependence on \(d_{\text{Cu-O}}\). It is noteworthy that the excitation energy exhibits such a systematic dependence on the local structure, and that it is apparently insensitive to whether the crystal has a planar, corner-sharing, or edge-sharing chain structure. This is consistent with our assignment of these features as MO excitations localized within a single Cu-O plaquette.

The overall trend exhibited in Fig. 2(a) is not unexpected. Intuitively, as the Cu and O atoms move closer, the \(p-d\) overlap will increase, and the Cu-O bonding becomes more covalent with a larger energy splitting \(\Delta_{\sigma^*}\). What is surprising is how strong this
\[ \Delta_{\sigma\sigma} \mathord{\sim} \Delta \mathord{\sim} d^0 \] and one then expects similar a \[ \sigma \] sion leads to \[ \Delta \mathord{\sim} d^0 \] states from the \[ \Delta \mathord{\sim} d^0 \] no dispersion of these features. The data of Fig. 2(b),
gest that the picture of a completely localized MO excita-
as a function of \[ p \mathord{\sim} eV \] is the hopping matrix element between the oxygen
expression.

Our RIXS measurements is different from the earlier re-
t

guides to the eye.

is data taken along the \((\pi \pi)\) left panel is along the \((\pi 0)\)

of \((2\pi 0)\) (\(\pi 0\)) (\(0 0\)) direction, and the right panel

\[ \Delta \mathord{\sim} d^0 \] dispersion becomes flat, as shown for the 2342 sample
(Figs. 2 and 3). Limited momentum dependence measure-
ments for the 2122 sample (not shown) also show a dispersionless behavior. The size of the observed disper-
sion along the Cu-O bond direction is plotted against \[ d_{\text{Cu-O}} \] in Fig. 2(b).

As shown in Fig. 2 the size of the dispersion and \[ \Delta_{\sigma\sigma} \mathord{\sim} d_{\text{Cu-O}} \] both increase as \[ d_{\text{Cu-O}} \] decreases. This suggests that the bandwidth of the dispersion is also controlled by the hopping
parameter \( t_{pd} \). Such behavior would, of course, be expected from the increased overlap of the wavefunctions,
since charge carriers then become less localized. How-
ever, the simplest picture of delocalized electrons fails
to describe the observed dispersion. For example, the
bandwidths of the Cu-O bonding and antibonding bands
in LCO are very large \( \sim 3 \) eV, and interband transitions
between these two bands would have a \textit{direct} gap of \( \sim 4 \) eV.

\section*{IV. DISCUSSION AND SUMMARY}

One of the most surprising results in this study is
that the \[ \Delta_{\sigma\sigma} \mathord{\sim} d_{\text{Cu-O}} \] scaling seems to apply to different
structures. In contrast, previous studies of the bond-length scaling of various quantities, such as charge-
transfer gap \( \Delta_{\text{CT}} \), or superexchange interaction \( J \),
have been limited to compounds withcorner-sharing
structures.\textsuperscript{20} It is well known that such quantities as superexchange coupling depend not only on the \( p-d \) hy-
bridization, but also crucially on the angle between the
two Cu-O bonds.\textsuperscript{23} One can argue that \[ \Delta_{\sigma\sigma} \] is a better
measure of Cu-O hybridization than \( J \) or \( \Delta_{\text{CT}} \), since it
is only dependent on \[ d_{\text{Cu-O}} \] and not on the presence of
neighboring atoms.

We have noted that the \( t_{pd} \sim d^{-8} \) dependence inferred from our study deviates significantly from the
tight-binding picture. It also appears to give rise to
discrepancies with other experiments. For example, for
materials with the corner-sharing structure, the three-
band Hubbard model gives \[ J \sim t_{pd}^2/\Delta_{\text{CT}}^2 \], where \( \Delta_{\text{CT}} \)
is an effective onsite Colomb interaction which is as-
sumed to be constant. If we use the experimentally
determined\textsuperscript{20} \[ \Delta_{\text{CT}} \sim d^{-6} \] and our \[ t_{pd} \sim d^{-8} \] result,
we obtain \( J \sim d^{-20} \), which clearly disagrees with the
much weaker \( d \)-dependence of \( J \) observed in various
experiments, including two magnon Raman scattering.\textsuperscript{3} One might expect that this discrepancy could be re-
solved by considering the fact that, due to strong elec-
tron correlations, the simple tight-binding picture of co-
valent bonding has to be modified. In fact, Mizuno
and coworkers\textsuperscript{3} considered two contributions to \( t_{pd} \).
That is, \( t_{pd} = t_{pd}^0 + t_{pd}^M \) where \( t_{pd}^0 \) is the contribution from the

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{dispersion.png}
\caption{The observed dispersion of the MO excitation. The left panel is along the \((\pi 0)\) direction, and the right panel is data taken along the \((\pi \pi)\) direction. The solid lines are guides to the eye.}
\end{figure}

\[ \Delta \mathord{\sim} d^0 \] dependence is. We have modeled the \[ d_{\text{Cu-O}} \] dependence of \[ \Delta_{\sigma\sigma} \] as a power law \( \Delta_{\sigma\sigma} \mathord{\sim} d^0 \) and
find \( \eta = -8(2) \), shown as a solid line in Fig. 2(a). Note
that the MO excitation energy is expected to be given
by
\[ \Delta_{\sigma\sigma} = \sqrt{(2t_{pp} - \Delta_0)^2 + 16t_{pd}^2}, \]
where \( t_{pp} \approx 0.65 \) eV is the hopping matrix element between the oxygen
\( p \) orbitals.\textsuperscript{29} Since \( 16t_{pd}^2 \gg (2t_{pp} - \Delta_0)^2 \), this
expression leads to \( \Delta_{\sigma\sigma} \approx 4t_{pd} \), to a first approximation,
and one then expects similar a \( d \)-dependence for \( \Delta_{\sigma\sigma} \) and \( t_{pd} \). Our results then implies that the \( d \)-dependence of \( t_{pd} \) is much stronger than that expected from tight-
binding theory, for which \( \sim d^{-3.5} \) is predicted.\textsuperscript{29} As plotted in Fig. 2(a), the observed RIXS data clearly devi-
ates from the \( \Delta_{\sigma\sigma} \approx d^{-3.5} \) behavior (dashed line). We
also note that the \( t_{pd} \sim d^{-8} \) behavior determined from our
RIXS measurements is different from the earlier re-
port of \( t_{pd} \sim d^{-4} \) by Cooper and coworkers, which was estimated indirectly from a three-band Hubbard model
expression.\textsuperscript{29}

In Fig. 2 the dispersion of the MO excitation is plotted
as a function of \( q \) for selected samples. These data sug-
uggest that the picture of a completely localized MO excita-
tion is an oversimplification – such a model would predict
no dispersion of these features. The data of Fig. 2(b),
which show significant dispersion for \( d_{\text{Cu-O}} < 1.93 \) Å,
suggest then that this localized picture breaks down for
small bond distances. For the LSCO17 sample, the
dispersion bandwidth is about 0.5 eV, with the minimum
excitation energy occuring at the zone boundary, imply-
ing an indirect gap. Note that this is completely different from the direct nature of the lower energy charge-transfer
gap as reported in Ref. 11. As \[ \Delta_{\sigma\sigma} \mathord{\sim} d_{\text{Cu-O}} \] decreases (e.g., LCO and LCCO), the dispersion of the MO excitation becomes
weaker. For the samples with an even smaller \[ \Delta_{\sigma\sigma} \mathord{\sim} d_{\text{Cu-O}} \] decreases. If we use the experimentally
determined \( \Delta_{\sigma\sigma} \mathord{\sim} d_{\text{Cu-O}} \) result, \( J \mathord{\sim} d^{-20} \),
which clearly disagrees with the
much weaker \( d \)-dependence of \( J \) observed in various
experiments, including two magnon Raman scattering.\textsuperscript{3} One might expect that this discrepancy could be re-
solved by considering the fact that, due to strong elec-
tron correlations, the simple tight-binding picture of co-
valent bonding has to be modified. In fact, Mizuno
atomic potential which depends only on $d_{\text{Cu-O}}$, while $t_{pd}^M$ is the contribution from the Madelung potential, which depends on the detailed arrangement of the neighboring ions. However, the calculated contribution from the Madelung potential is of order of $\sim 0.1$ eV or smaller, so that this alone is not enough to explain the $\sim d^{-8}$ dependence.

These results may be suggesting that one has to abandon the simple relationship of $\Delta_{\sigma\sigma^*} \approx 4t_{pd}$. Certainly, as discussed above, the picture of a completely localized MO excitation is apparently an oversimplification, since it breaks down as $d_{\text{Cu-O}}$ becomes shorter – as evidenced by the sizable dispersion observed in LSCO17. Thus, if a more realistic expression for $\Delta_{\sigma\sigma^*}$ is used, $t_{pd} \sim d^{-3.5}$ scaling law might be recovered. For example, a recent first-principles calculation has emphasized the role of apical oxygens in the systematics of high temperature superconductivity. Indeed, the scaling plot in Fig. 2 also exhibits some systematic dependence on the number of apical oxygens and it may be interesting to further investigate the role played by apical oxygens. Certainly, a systematic ab initio calculation of the $d_{\text{Cu-O}}$-dependence of MO excitation energy in large clusters would be highly desirable and may help to clarify the relationship between $\Delta_{\sigma\sigma^*}$, $t_{pd}$, and $d_{\text{Cu-O}}$.

To summarize, we have studied a charge-transfer excitation in various cuprate compounds using resonant inelastic x-ray scattering technique. We assign this excitation to a mostly localized molecular orbital excitation, that is, an excitation from a bonding to an antibonding molecular orbital. We have found that this molecular orbital excitation energy, which is a measure of the hopping matrix element $t_{pd}$, exhibits a systematic Cu-O bond length dependence, which is much stronger than that expected from tight-binding theory. We have also observed a sizable dispersion of this excitation in some materials, suggesting that this molecular orbital excitation becomes less localized as the $p-d$ hybridization becomes large.

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35 There are two apical oxygens in the structure of LSCO17, LSCO5, and LCO, while there is one apical oxygen for LCCO. The rest of the samples have no apical oxygens.