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Optical determination of the relation between the electron-boson coupling function and the critical temperature in high-\(T_c\) cuprates

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We take advantage of the connection between the free-carrier optical conductivity and the glue function in the normal state, to reconstruct from the infrared optical conductivity the glue spectrum of ten different high-\(T_c\) cuprates revealing a robust peak in the 50–60 meV range and a broad continuum at higher energies for all measured charge-carrier concentrations and temperatures up to 290 K. We observe that the strong-coupling formalism accounts fully for the known strong temperature dependence of the optical spectra of the high-\(T_c\) cuprates, except for strongly underdoped samples. We observe a correlation between the doping trend of the experimental glue spectra and the critical temperature. The data obtained on the overdoped side of the phase diagram conclusively exclude the electron-phonon coupling as the main source of superconducting pairing.

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I. INTRODUCTION

The theoretical approaches to the high-\(T_c\) pairing mechanism in the cuprates are divided in two main groups: According to the first electrons form pairs due to a retarded attractive interaction mediated by virtual bosonic excitations in the solid. These bosons can be lattice vibrations, fluctuations of spin polarization, electric polarization, or charge density. The second group of theories concentrates on a pairing mechanism entirely due to the nonretarded Coulomb interaction or so-called Mottness. Indeed, optical experiments have found indications for mixing of high and low energy degrees of freedom when the sample enters into the superconducting state.

An indication that both mechanisms are present was obtained by Maier, Poilblanc, and Scalapino, who showed that the “anomalous” self-energy associated with the pairing has a small but finite contribution extending to an energy as high as \(U\), demonstrating that the pairing interaction is, in part, nonretarded. The experimental search for a pairing glue will play an essential role in determining the origin of the pairing interaction. Aforementioned glue is expressed as a spectral density of these bosons indicated as \(\Phi^\text{pair}(\omega)\) for phonons and \(\hat{I}^2\chi(\omega)\) for spin fluctuations, here represented as the general, dimensionless function \(\hat{\Pi}(\omega)\). An important consequence of the electron-boson coupling is that the energy of the quasiparticles relative to the Fermi level \(\xi\) is renormalized and their lifetime becomes limited by inelastic decay processes involving the emission of bosons. The corresponding energy shift and the inverse lifetime, i.e., the real and imaginary parts of the self-energy are expressed as the convolution of the “glue function” \(\hat{\Pi}(\omega)\) with a kernel \(\hat{K}(\xi,\omega,T)\) describing the thermal excitations of the glue and the electrons.

\[
\Sigma(\xi) = \int K(\xi,\omega,T)\hat{\Pi}(\omega)d\omega.
\]

In the absence of a glue and of scattering off impurities the effect of applying an ac electric field to the electron gas is to induce a purely reactive current response, characterized by the imaginary optical conductivity \(\Im[\sigma(\omega)] = \omega^2 / \omega + M(\omega)\), where the plasma frequency, \(\omega_p\), is given by the (partial) f-sum rule for the conduction electrons. The effect of coupling the electrons to bosonic excitations is revealed by a finite, frequency dependent dissipation, which can be understood as arising from processes whereby a photon is absorbed by the simultaneous creation of an electron-hole pair and a boson. As a result, the expression for the optical conductivity in the normal state,

\[
4\pi\sigma(\omega) = \frac{i\omega^2}{\omega + M(\omega)},
\]

now contains a memory function or optical self-energy. A particularly useful aspect of this representation is that \(\hat{M}(\omega)\) follows in a straightforward way from the experimental optical conductivity. The optical self-energy is related to the single-particle self-energies by the expression

\[
\frac{\hat{M}(\omega)}{\omega} = \left\{ \int \frac{f(\xi) - f(\xi + \omega)}{\omega + \Sigma'(\xi) - \Sigma(\xi + \omega)}d\xi \right\}^{-1} - 1.
\]

The central assumption in the above is the validity of the Landau Fermi-liquid picture for the normal state. The aforementioned strong-coupling analysis is therefore expected to work best on the overdoped side of the cuprate phase diagram, where the state of matter appears to become increasingly Fermi liquidlike. If antiferromagnetism is necessary to obtain the insulating state in the undoped parent compounds, as has been argued based on the doping trends of the Drude spectral weight, the strong-coupling analysis may in prin-
ciple be relevant for the entire doping range studied. However, in the limit of strong interactions aforementioned formalism needs to be extended, e.g., with vertex corrections and it eventually breaks down. We therefore define the function $\tilde{\Pi}(\omega)$ as the effective quantity which in combination with Eqs. (1) and (3), returns the exact value of $M(\omega)$ for each frequency. Defined in this way $\tilde{\Pi}(\omega)$ captures all correlation effects regardless of whether the system is a Fermi liquid or not. This becomes increasingly relevant when the doping is lowered below optimal doping.

Here we take advantage of the connection between the temperature and frequency-dependent conductivity in the normal state and the glue spectrum to test experimentally the consequences of the standard approach to check the internal consistency of it and to determine the range of doping where internal consistency is obtained. For a $d$-wave superconductor, the momentum dependence is essential to understand the details of the pairing. This, of course, is difficult to handle for optical spectroscopy which is inherently a momentum integrated probe. Nevertheless, optical spectra provide the important information on the energy scale of the bosons involved and on the doping and temperature evolution. The paper is organized as follows. In Sec. II we show that the temperature dependence of the optical spectra of the cuprates is well described within the strong-coupling formalism described above. In Sec. III we present the $\tilde{\Pi}(\omega)$ functions for ten different cuprates. These $\tilde{\Pi}(\omega)$ functions are used in Sec. IV to estimate critical temperatures and Sec. V discusses the implications of these results with regard to the pairing mechanism in the cuprates. Finally, in Sec. VI we summarize our results.

II. INTERNAL CONSISTENCY CHECK OF THE STRONG-COUPLING FORMALISM

In order to test whether the strong-coupling analysis is applicable to the cuprates we start with an important test of its internal consistency: (i) we invert the data at 290 K to obtain $\tilde{\Pi}(\omega)$, (ii) we use this $\tilde{\Pi}(\omega,290)$ K to predict the optical spectra at lower temperatures. If the prediction faithfully reproduces the experimental spectra at these temperatures, we have a strong indication that the electronic structure and its evolution as a function of temperature are to a good approximation within the realm of strong-coupling theory. We use a standard least-squares routine to fit a histogram representation of $\tilde{\Pi}(\omega)$ to our experimental infrared spectra (see the Appendix). The quantity $\tilde{\Pi}(\omega)$ is shown in Fig. 1 for optimally doped HgBa$_2$CuO$_{4+\delta}$(Hg-1201) (Ref. 18) for $T=290$ K together with the optical self-energies calculated from this function at three different temperatures. For 290 K the theoretical curve runs through the data points, reflecting the full convergence of the numerical fitting routine.

It is interesting to notice that the shoulder at 80 meV in the 100 K experimental data is reproduced by the same $\tilde{\Pi}(\omega)$ function as the one used to fit the 290 K data. It can be excluded that this shoulder is due to the pseudogap since a gap is certainly absent for temperatures as high as 290 K. The shoulder is therefore entirely due to coupling of the electrons to a mode at approximately 60 meV. On the other hand, the considerable sharpening of this feature for temperatures lower than 100 K finds a natural explanation in the opening of a gap, as illustrated in the inset of Fig. 1. We see that for 100 K the theoretical prediction also runs through the experimental data points. In other words, the strong temperature dependence of the experimental optical spectra is entirely due to the Fermi and Bose factors of Eqs. (1) and (3).

This example confirms the close correspondence between the features in $M(\omega)$ and in $\tilde{\Pi}(\omega)$ pointed out in Ref. 19. In particular the broad maximum in $M(\omega)$ has its counterpart in the high-intensity region of $\tilde{\Pi}(\omega)$ terminating at 290 meV. The internal consistency is therefore demonstrated by the fact that the large temperature dependence of the optical spectra is fully explained by the strong-coupling formalism. This consistency was obtained for all samples, except for the most strongly underdoped single-layer Bi2201 sample.

III. ELECTRON-BOSON COUPLING FUNCTION

As summarized in Fig. 2, we have analyzed previously published optical spectra of ten different samples belonging to different families of materials, i.e., optimally doped Hg-1201 (Ref. 18) and Bi$_2$Sr$_2$Ca$_2$Cu$_2$O$_{8+\delta}$ (Bi-2223), as well as four Bi$_2$Sr$_2$Ca$_2$Cu$_2$O$_{8+\delta}$ (Bi-2212) crystals with different hole concentrations. In addition, we analyzed data for four Bi$_2$Sr$_2$Ca$_2$Cu$_2$O$_{8+\delta}$ (Bi-2201) crystals with different hole concentrations.

Excellent fits were obtained for all temperatures but the $\tilde{\Pi}(\omega)$ spectra exhibit a significant temperature dependence in particular at the low-frequency side of the $\tilde{\Pi}(\omega)$ spectrum. Since all thermal factors contained in Eqs. (1) and (3) are, in principle, folded out by our procedure, the remaining temperature dependence of $\tilde{\Pi}(\omega)$ reflects the thermal properties of the “glue function” itself. Such temperature dependence is a direct consequence of the peculiar dc and far-infrared conductivity, in particular the $T$-linear dc resistivity and $\omega/T$ scaling of $\sigma(\omega,T)$ at optimal doping. For the highest doping levels both $\tilde{\Pi}(\omega)$ and its temperature dependence diminish, which is an indication that a Fermi-liquid regime is approached. The most strongly underdoped sample Bi-2201-UD0 exhibits an upturn of the imaginary part of the experimental optical self-energy for $\omega \rightarrow 0$. This aspect of the data cannot be reproduced by the strong-coupling expression, resulting in an artificial and unphysical peak at $\omega=0$ of the fitted $\tilde{\Pi}(\omega)$ function.

We observe two main features in the glue function: A robust peak at 50–60 meV and a broad continuum. The upper limit of $\tilde{\Pi}(\omega)$ is situated around approximately 300 meV for optimally doped single-layer Hg1201 and for the bilayer and trilayer samples. The continuum extends to the highest energies (550 meV for the single-layer samples and 400 meV for the bilayer) for the weakly overdoped samples, whereas the continuum of the strongly doped bilayer sample extends to only 300 meV. This high energy scale of the glue function...
The most prominent feature, present in all spectra reproduced in Fig. 2 is a peak corresponding to an average frequency of $60 \pm 3$ meV at room temperature (see the Appendix for an estimate of the error bar). Perhaps the most striking aspect of this peak is the fact that its energy is practically independent of temperature (up to room temperature) and sample composition. Moreover, the intensity and width are essentially temperature independent. While our results confirm by and large the observations of Hwang et al.,23,24 in the pseudogap phase, the persistence of the 50–60 meV peak to room temperature has not been reported before for these compounds. However, Collins et al. obtained excellent fits to their infrared data of YBa$_2$Cu$_3$O$_7$ at 100 and 250K using for both temperatures the same $\alpha^2 F(\omega)$ spectrum with a peak at $\sim 35$ meV and a continuum extending up to 300 meV. The 50–60 meV peak which we observe, arises most likely from the same boson that is responsible for the “kink” seen in angle resolved photoemission (ARPES) experiments along the nodal direction in $k$ space at approximately the same energy.25–27 The peak-dip-hump structure in the tunneling spectra (STS) (Refs. 28–30) has also been reported at approximately the same energy.

IV. CRITICAL TEMPERATURES

One of the most important issues in the field of high $T_c$ is the question whether pairing is caused by the exchange of virtual bosons. These processes are described by a bosonic density-of-states function closely related to $\Pi(\omega)$. If the electron-electron interaction occurs uniquely in the $d$-wave channel, the superconducting critical temperature follows from the usual relation

$$T_c = 0.83 \bar{\omega} \exp\left(-\frac{1}{1 + \lambda_d/\lambda_d}\right), \tag{4}$$

where $\lambda_d$ is the coupling constant in the $d$-wave pairing channel and

$$\lambda_d = 2 \int_0^{\infty} \Pi_d(\omega)/\omega d\omega, \tag{5}$$

and $\Pi_d(\omega)$ is the $d$-wave electron-boson coupling function. The effective frequency of the bosons responsible for the pairing interaction is obtained by taking the average of $\ln(\omega)$ weighted by electron-boson coupling function,31

$$\ln(\bar{\omega}) = 2 \lambda^{-1} \int_0^{\infty} \omega^{-1} \Pi_d(\omega) \ln(\omega) d\omega. \tag{6}$$

To apply Eq. (4) to our experimentally measured $\Pi(\omega)$, we would need to map this function on the $d$-wave pairing channel. Boson fluctuations below a certain critical frequency act as pair breakers, as has been shown by Millis, Varma, and Sachdev in the case of spin-fluctuation-mediated $d$-wave superconductivity. Clearly, it is not possible to separate pair-breaking from pair-forming contributions to $\Pi(\omega)$ in an unambiguous way. To proceed we assume that the full $\Pi(\omega)$ function contributes favorably to the pairing. This means that our results overestimate the critical temperature. In Table I...
we indicate the total coupling constant $\lambda$ and logarithmic frequency $\tilde{\omega}$ for the room-temperature $\tilde{\Pi}(\omega)$ spectra. The coupling strength shows a strong and systematic increase with decreasing hole concentration, which probably requires a theoretical treatment beyond the strong-coupling expansion. At the same time we see that $\tilde{\omega}$ shows the opposite trend.

An estimate of $T_c$, using the experimental values indicated in Table I gives values in the $100–200$ K range. The critical temperature can also be calculated straightforwardly from the $s$-wave Eliashberg equations when $\tilde{\Pi}(\omega)$ is known. As shown in Fig. 3, the $T_c$ are in the $150–300$ K range and they correlate with the experimentally observed doping trends of $T_c$. The dome-shaped trend in the calculation is a consequence of the increasing energy scale of $\tilde{\Pi}(\omega)$ and the decreasing overall coupling constant as a function of doping.

V. IMPLICATIONS FOR THE PAIRING MECHANISM

We take this analysis a step further by calculating $T_c$ from the optical spectra below $100$ meV ($\tilde{\Pi}_{pk}$) and above $100$ meV ($\tilde{\Pi}_{cnt}$). The resulting coupling constants and $T_c$s are indicated in Table I. On the underdoped side $\tilde{\Pi}_{cnt}$ vanishes, and $T_c$ is given only by the coupling to the intense $50–60$ meV peak in $\tilde{\Pi}(\omega)$, but in this doping range we have to be careful with the interpretation of our results. As mentioned in the introduction our $\tilde{\Pi}(\omega)$ spectra represent effective coupling functions, which may contain effects arising from features not captured by the strong coupling Eqs. (1) and (3). If, for example, a pseudogap opens in the electronic spectrum this will affect the shape of $\tilde{\Pi}(\omega)$. These effects likely play a role for the underdoped samples, but are not expected to affect much the room-temperature values, indicated in Table I and Fig. 3. On the contrary, the larger temperature dependence seen for underdoped samples in Fig. 2 may well be a result of the opening of a pseudogap. For the overdoped samples the $T_c$s calculated from $\tilde{\Pi}_{pk}$ are smaller than the experimental values. For example, for Bi2212 with the highest doping $\tilde{\Pi}_{pk}$ gives only $T_c<20$ K, whereas $\tilde{\Pi}_{cnt}$ gives $160$ K, implying that the glue function above $100$ meV is of crucial importance for the pairing mechanism. Since only electronic modes can have such high energies, an important contribution to the high $T_c$ mechanism comes apparently from coupling to electronic degrees of freedom, i.e., spin$^{13,12,19}$ or orbital current fluctuations.$^2$

VI. SUMMARY

In summary, the $\tilde{\Pi}(\omega)$ spectrum obtained from the optical spectra of ten different compounds using a strong-coupling analysis is observed to consist of two features: (i) a robust peak in the range of $50–60$ meV and (ii) a doping dependent continuum extending to $0.3$ eV for the samples with the highest $T_c$. We perform an important test of the internal consistency of the strong-coupling formalism by showing that the temperature dependence of the optical spectra is determined by Fermi and Bose factors in the strong-coupling expressions. The remaining temperature dependence of $\tilde{\Pi}(\omega)$ can therefore be taken to indicate that part of the $\tilde{\Pi}(\omega)$ spectrum is electronic in origin. We observe an intriguing corre-
lation between the doping trend of the experimental glue spectra and the critical temperature. Finally we obtain an upper limit to the contribution of electron-phonon coupling to the pairing of the overdoped samples, which is too small to account for the observed critical temperature.

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APPENDIX

The inversion of Eqs. (1) and (2) allows to extract $\tilde{\Pi}(\omega)$ from experimental data of the optical conductivity or related optical spectra. The accuracy of the resulting $\tilde{\Pi}(\omega)$ spectrum is in practice limited by the convolution with thermal factors expressed by Eqs. (1) and (2). Microscopic models giving roughly the same $\tilde{\Pi}(\omega)$ spectra, which differ however in the details of the frequency dependence of this quantity may therefore provide fits to the directly measured optical quantities such as infrared reflectance spectra, which at first glance look satisfactory but the remaining discrepancies with the experimental spectra may nevertheless be of significant importance for the proper understanding of the optical data. It is therefore of crucial importance to test the “robustness” of each fit with regard to the spectral shape of the $\tilde{\Pi}(\omega)$ function imposed by such models. This robustness can be tested by including in the fit routine one or several “oscillators” superimposed on the model function. When the model glue function provides a complete description of the electronic structure, adding extra oscillators will not result in an improvement of the quality of the fit. We have used this approach to test functional forms commonly used in the literature, in particular the marginal Fermi-liquid (MFL) model and the Millis-Monien-Pines (MMP) representation of the spin-fluctuation spectrum.

We found that neither of these functional forms describe completely the experimental data. In search of a more flexible form of $\tilde{\Pi}(\omega)$ we used a superposition of lorentzian oscillators and found that it could be used to describe all available experimental data in a consistent manner. The resulting $\tilde{\Pi}(\omega)$ functions and trends are equivalent to those in Fig. 2. From these initial tests we concluded that due to the thermal smearing expressed by Eqs. (1) and (2) our $\tilde{\Pi}(\omega)$ spectra can only be determined with limited resolution. This lead us to the use of a histogram representation, where each block in the histogram represents a likelihood to find coupling to a mode with a well-determined coupling strength. For the lowest frequency interval ($0<\omega<\omega_1$) a triangular shape was used instead of a block, which is necessary to avoid problems with the convergence of the integral $\int_0^{\omega_1} \omega^{-1} \tilde{\Pi}(\omega) d\omega$. In practice the output generated by the fitting routine has low intensity in this first interval and the triangles are therefore difficult to distinguish in Fig. 2.

To give an example: the block centered at 55 meV seen in the Hg-1201 sample in Fig. 2 has $\lambda=1$ and a width of about 30 meV. Our histogram representation implies the presence of a coupling to one or several modes between 45 and 75 meV with an integrated coupling strength of 1. The histograms thus constitute the most detailed representation of $\tilde{\Pi}(\omega)$ given the precision of our experimental reflectivity and ellipsometry spectra.

Examples of experimental reflectivity data together with the fits are shown in Fig. 4 for a selection of representative samples and temperatures. (a): underdoped nonsuperconducting Bi-2201 (UD0). (b): optimally doped Hg-1201 with $T_c = 97$ K (OpD97) (Ref. 18) (c): overdoped nonsuperconducting Bi-2201 (OD0). Weak sharp peaks, particularly visible for the strongly underdoped sample in panel (a) are due to transverse optical phonons, which we do not intend to fit.

\[
=2\int_0^{\omega_1} \omega^{-1} \tilde{\Pi}(\omega) d\omega. \]

FIG. 4. (Color) Experimental reflectivity (red and blue lines) and fit curves (black lines) for selected samples and temperatures. (a): underdoped nonsuperconducting Bi-2201 (UD0). (b): optimally doped Hg-1201 with $T_c = 97$ K (OpD97) (Ref. 18) (c): overdoped nonsuperconducting Bi-2201 (OD0). Weak sharp peaks, particularly visible for the strongly underdoped sample in panel (a) are due to transverse optical phonons, which we do not intend to fit.
When $N$ is summed [see Eq. (1)] is $N=400$. The quoted values of $\chi^2$ are those for the room-temperature spectra. At 100 K these values increase by a factor 1.5 (MFL) and 3.5 (MMP). In contrast, in the oscillator and the histogram model the $\chi^2$ was found to be independent of temperature.

$$\chi^2 \equiv \sum_{i=1}^{N} \left[ \frac{R(\omega_i) - f(\omega_i;p_1, \ldots, p_n)}{\sigma_i} \right]^2,$$

where $R(\omega)$ is an experimentally measured data point, $f(\omega_1,p_1, \ldots, p_n)$ is the calculated value in this point based on parameters $p_1, \ldots, p_n$, and the difference between these two is weighed by the error bar $\sigma_\omega$ determined for $R(\omega)$. For a given set of reflectivity and ellipsometry data at one particular temperature, using a standard PC, the iteration takes about 3 h until convergence is reached. The Levenberg-Marquardt least-squares method is an extremely powerful method to find the minimum of $\chi^2$ in a multidimensional parameter space. To ensure that $\chi^2$ has converged to the global minimum in parameter space several tests have been performed, for each individual sample and temperature displayed in Fig. 5, where in each test the optimization process was started from a different set of starting parameters. To give some idea of the robustness of our method we will here discuss one representative example: optimally doped Hg-1201.

The models are evaluated based on the minimum found for $\chi^2$. A comparison of Figs. 5(a)–5(d) shows that the MMP model describes better the optical data then the MFL model but that they give similar results if we add an extra oscillator to these models. Figures 5(e) and 5(f) show the model independent results mentioned above and are very similar to the modified MMP and MFL model. The models in these last two panels have the same $\chi^2$ and the comparison in [Fig. 5(g)] shows that the histogram representation realistically expresses the uncertainty in the position of the low energy peak, while the correspondence between the features in both models remains excellent. It is interesting that the model with two oscillators is described by six parameters, while the histogram representation uses 12 parameters. The fact that the fit routine adjusts the latter 12 parameters in such a manner as to reproduce the two oscillators, proves that the features represented in the right-hand panel of Fig. 5 are realistic.

The models presented in Fig. 5 allow us to make an estimate of the uncertainty in the determination of the frequency of the low energy peak. We define the first moment of this peak as

$$\langle \omega \rangle = \int_0^{100 \text{ meV}} \omega \tilde{\Pi}(\omega) d\omega / \int_0^{100 \text{ meV}} \tilde{\Pi}(\omega) d\omega.$$

The variance of $\langle \omega \rangle$ is defined as

$$\sigma^2 = \frac{1}{N} \sum_{i=1}^{N} ((\omega_i) - \langle \omega \rangle)^2$$

with $\langle \omega \rangle$ the mean of the moments of the spectra presented in Fig. 5 and $i=1 \ldots N$ runs over the number of spectra used ($N=6$ for each temperature). For the Hg1201 room-temperature spectra presented in Figs. 5(a)–5(f) we find $\langle \omega \rangle \approx 60$ meV and $\sigma \approx 3$ meV. This value is approximately the same for all samples. In Fig. 6 we present the temperature dependence of the first moment of the glue functions presented in Fig. 2.
OPTICAL DETERMINATION OF THE RELATION BETWEEN…

13 $\frac{\beta}{\gamma} = \int \frac{n(\omega + \epsilon) - n(\omega)}{\epsilon} d\epsilon$ where $n(\omega)$ and $f(\epsilon)$ are the Bose and Fermi-Dirac distribution functions respectively (Ref. 16).
15 An alternative frequently used notation for this memory function (Ref. 23) is $-2\Delta\tilde{\gamma}(\omega) = \tilde{M}(\omega)$.