Optical study of superconducting Pr$_2$CuO$_x$ with $x \simeq 4$

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Superconducting Pr$_2$CuO$_x$, $x \simeq 4$ (PCO) films with $T'$ structure and a $T_c$ of 27 K have been investigated by various optical methods in a wide frequency ($7 - 55000$ cm$^{-1}$) and temperature (2 to 300 K) range. The optical spectra do not reveal any indication of a normal-state gap formation. A Drude-like peak centered at zero frequency dominates the optical conductivity below 150 K. At higher temperatures, it shifts to finite frequencies. The detailed analysis of the low-frequency conductivity reveals that the Drude peak and a far-infrared (FIR) peak centered at about 300 cm$^{-1}$ persists at all temperatures. The FIR-peak spectral weight is found to grow at the expense of the Drude spectral weight with increasing temperature. The temperature dependence of the penetration depth follows a behavior typical for $d$-wave superconductors. The absolute value of the penetration depth for zero temperature is 1.6 µm, indicating a rather low density of the superconducting condensate.

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I. INTRODUCTION

It is commonly accepted that the parent compounds of the superconducting high-$T_c$ cuprates are antiferromagnetic charge-transfer insulators and that superconductivity emerges upon doping either with holes or electrons. There are some similarities but also differences between hole- and electron-doped cuprates. One similarity is that all cuprate superconductors have a perovskite structure with the common feature of square planar copper-oxygen planes separated by rare-earth oxide (charge-reservoir) layers. On the other hand, they differ in that the hole- and electron-doped cuprates. One similarity is that all cuprate superconductors have a perovskite structure with the common feature of square planar copper-oxygen planes separated by rare-earth oxide (charge-reservoir) layers.

There are also a large difference between the phase diagrams of hole- and electron-doped cuprates. Whereas the antiferromagnetic phase exists only over a small doping range ($0 - 4\%$) in hole-doped cuprates, it is more robust in electron-doped cuprates and persists to higher doping levels ($0 - 11\%$). Superconductivity occurs in a doping range that is almost five times narrower for electron-doped cuprates ($11 - 17\%$) as compared to the hole-doped counterparts ($4 - 32\%$). While consensus on the phase diagram exists for the hole-doped side, the situation for the electron-doped cuprates is less obvious.

As early as in 1995, Brinkmann et al. demonstrated that the superconductivity window in Pr$_{2-x}$Ce$_x$CuO$_4$ single crystals can be extended down to a doping level of 4% by a special oxygen reduction and annealing technique. Improved deposition and annealing techniques have recently made it possible to produce thin films of electron-doped parent compounds (R$_2$CuO$_4$, $R$ = Pr, Sm, Nd, Eu, and Gd) with $T'$ structure that, in fact, are metallic and superconducting at low temperatures.

This sharp contradiction to earlier results is explained as being due to the fact that although apical oxygen should not exist in the ideal $T'$ structure, in practice (especially in bulk samples) it is usually not completely removed. This apical oxygen in the $T'$ structure acts as a very strong scatterer and pair breaker. In contrast to bulk samples, the large surface-to-volume ratio of thin films along with their tenuity itself is advantageous in achieving the proper $T'$ structure with no apical oxygen.

The reported superconductivity in undoped cuprates puts a question mark on the applicability of the charge-transfer-insulator picture to electron-doped cuprates. Remarkably, recent calculations on the basis of a newly developed first-principles method show a radical difference between the parent compounds with $T$ and $T'$ structures. The first are found to be charge-transfer insulators, while the latter, e.g., Pr$_2$CuO$_4$, are essentially metallic and their apparent insulating nature may originate from magnetic long-range order (Slater transition).
which is competing with the metallic ground state.\textsuperscript{16} One should note, however, that it is still a question whether or not $T'$ superconductors are truly undoped or are still doped by possible oxygen vacancies in the RO layers during the reduction process. Since bulk $T'$-$R_2$CuO$_4$ superconducting samples have not yet been synthesized, direct measurements of the oxygen distribution are not available so far. Nevertheless, neutron diffraction on Nd$_{2-x}$Ce$_x$CuO$_{4+y}$ single crystals shows that it is mostly apical oxygen which is removed during reduction.\textsuperscript{17,18} The synthesis of bulk samples of a nominally undoped $T'$-(La,Sm)$_2$CuO$_4$\textsuperscript{19,20} and of heavily underdoped Pr$_{1.3-x}$La$_{0.7}$Ce$_x$CuO$_{4+y}$\textsuperscript{21} gives hope that the oxygen stoichiometry might be determined in the near future for this class of superconductors.

In this paper, we do not touch the issue of oxygen stoichiometry; instead we present a comprehensive broadband optical investigation of Pr$_2$CuO$_y$ (PCO) films with $x \approx 4$. As argued above, it is impossible to rule out doping by oxygen vacancies (if this is the case, $x$ differs from 4 in our films). However, we will show that our findings can also be consistently understood within the picture, where superconductivity develops in undoped PCO (i.e., $x = 4$). We demonstrate that the available PCO samples do show a metallic as well as a superconducting optical response. We find that this response can be reconciled with $d$-wave superconductivity and the density of the superconducting condensate is rather low. We do not observe any indication of a normal-state pseudogap. All this supports ideas that the standard charge-transfer-insulator picture might not be applicable to PCO.

## II. EXPERIMENT

PCO films were grown by molecular beam epitaxy (MBE)\textsuperscript{22} on a (110)-oriented 0.35 mm thick DyScO$_3$ substrate. The phase purity of these films was confirmed by x-ray diffraction. The films were 100 nm thick with the $c$ axis oriented perpendicular to the film’s surface. Direct-current (dc) resistivity was measured from 4 to 300 K by a standard four-probe method.

Near-normal reflectivity from 40 to 55000 cm$^{-1}$ (5 − 6800 meV) was measured using a combination of two Fourier-transform spectrometers (Bruker IFS113V and Bruker IFS86V/s) covering frequencies from 40 to 22000 cm$^{-1}$ and a grating spectrometer for room-temperature reflectivity measurements from 8000 to 55000 cm$^{-1}$. In order to obtain the absolute reflectivity of the sample, we used an in situ gold (for the infrared) or silver (for the visible) overfilling technique.\textsuperscript{22} With this technique, we achieved an absolute accuracy in the reflectivity better than 3 % and the relative error between different temperatures was of the order of 0.5 %. The room temperature reflectivity in the ultraviolet was measured against an aluminum mirror and then corrected for the absolute reflectivity of aluminum.

Normal-incident phase-sensitive transmission at 210 and 250 GHz (7 and 8.3 cm$^{-1}$) was measured as a function of temperature with a spectrometer employing backward-wave oscillators (BWOs) as sources of coherent radiation.\textsuperscript{22} A Mach-Zehnder interferometer arrangement of the spectrometer allows measurements of both the intensity and the phase shift of the wave transmitted through the sample. Using the Fresnel optical formulas for the complex transmission coefficient of the two-layer system, the film’s complex conductivity as well as the penetration depth were directly obtained from these measurements. This experimental method has been previously applied to a large number of different superconductors.\textsuperscript{22} Technical details of our experimental procedure can be found in Ref.\textsuperscript{22}

Optical properties of bare substrates were obtained from measurements performed in the same frequency and temperature windows as for the thin-film samples.

We investigated two thin films of PCO. The results, obtained on the films, do not demonstrate any significant difference. Hereafter we present results for one of the two films.

## III. RESISTIVITY

Fig. 1 shows the temperature dependence of the resistivity of the PCO film. The resistivity decreases monotonically with decreasing temperature down to $T_c = 27$ K. The width of the superconducting transition is 0.8 K. The temperature dependence of the resistivity can be described by the power law

$$\rho(T) = \rho_0 + AT^n,$$

with $\rho_0 = 0.151$ m$\Omega$cm, $A = 10^{-5}$ m$\Omega$cmK$^{-n}$, and $n = 2$ from $T_c$ up to 150 K. The quadratic temperature dependence is in agreement with earlier reports on superconducting Nd$_{2-x}$Ce$_x$CuO$_4$ films and single crystals for tem-
IV. OPTICAL PROPERTIES

A. Raw experimental data

Figure 2 shows the as-measured in-plane (ab-plane) reflectivity of the PCO film on a DyScO$_3$ substrate versus frequency at various temperatures. At low frequencies, the reflectivity is quite high and increases with decreasing temperature, typical for metals. A number of phonon modes from the substrate and the film appear at frequencies below 700 cm$^{-1}$. The maxima seen above some 10000 cm$^{-1}$ can be attributed to interband transitions.

The changes to the reflectivity spectra induced by the superconducting transition are not very well pronounced within our experimental accuracy. This is because of a relatively high transparency of the film. Thus, the results, obtained from the reflectivity measurements, are only discussed in the normal state in the course of the article.

The formation of the superconducting condensate can instead be directly seen by use of our low-frequency phase-sensitive transmission measurements. In Fig. 3 we present examples of these measurements. The onset of the transition into the superconducting state reveals itself immediately as a reduction of the temperature-dependent power transmission $T_r$ and the phase shift. However, the penetration depth and the superfluid density, obtained from these measurements, are discussed in Sec. IV.B.

B. Normal-state optical conductivity

By applying a thin-film fitting procedure, described in detail in App. A, we extract the film’s complex optical conductivity, $\sigma = \sigma_1 + i\sigma_2$, from our reflectivity spectra. Neither BWO data nor values of the dc conductivity in the normal state have been utilized within this fitting procedure.

The real part of the PCO optical conductivity obtained by this modeling is shown in Fig. 4 for various temperatures indicated in the legend. As the lowest frequency of the reflectivity measurements was 40 cm$^{-1}$ the data obtained from this analysis below this threshold frequency are to be considered as extrapolations and, thus, are shown as dashed lines. Nevertheless, the zero-frequency limit of $\sigma_1$ evolves in accordance with $\sigma_{dc}$ at all temperatures in the normal state (bold points on the vertical left-hand axis of Fig. 4).

At all temperatures above $T_c$, the optical conductivity of PCO can be disentangled into a Drude component and a set of Lorentz oscillators, representing a broad far-infrared (FIR) band, narrow FIR peaks, a mid-infrared (MIR) band, and interband-transition bands at the high-

FIG. 2: (Color online) Reflectivity of the PCO thin film on a DyScO$_3$ substrate as a function of frequency at various temperatures listed in the legend. The $E$ vector of the probing radiation lies in the $ab$ plane of the film (and parallel to the [001] axis of the substrate). The inset shows the reflectivity of the bare substrate at 4 K.

FIG. 3: (Color online) Examples of raw (i.e. not normalized to the empty-channel measurements) phase-sensitive transmission measurements at 8.3 cm$^{-1}$. Power transmission $T_r$ (top panel) and phase shift (middle panel) of the wave passed through the PCO film on the DyScO$_3$ substrate are shown as a function of temperature together with a close-up of the dc resistivity measurements. The thin vertical line indicates $T_c$. Power transmission $T_r$ (bottom panel) and phase shift (bottom panel) of the wave passed through the PCO film on the DyScO$_3$ substrate are shown as a function of temperature together with a close-up of the dc resistivity measurements. The thin vertical line indicates $T_c$. The pene-
The authors have elaborated on the possible origin of these additional modes but concluded eventually that some impurities and/or contributions from different phases may play a role. This conclusion is perfectly in line with claims made in Refs. 4–10 that the complete removal of all apical oxygen is extremely challenging and absolutely necessary for superconductivity in undoped T' cuprates.

The bump at about 300 cm\(^{-1}\) in the \(\sigma_1(\omega)\) spectra can be attributed to electron localization. Such behavior is known for the superconducting cuprates\(^{27}\) and is typical for so-called bad metals where a certain degree of disorder is inherently present.\(^{33,34}\)

Using dynamical mean-field theory (DMFT) and iterated perturbation theory, Mutou and Kontani demonstrated\(^{24}\) that in a strongly correlated metallic state, realized by a large on-site repulsion energy, the optical conductivity develops a Drude peak centered at \(\omega = 0\) at low temperatures and a shift of this peak to finite frequencies above the Ioffe-Regel-limit temperature \(T_{IR}\), although the resistivity increases monotonically even at \(T > T_{IR}\).

A temperature evolution of far-infrared \(\sigma_1(\omega)\), similar to the one observed here, has been reported for underdoped Nd\(_{2-x}\)Ce\(_x\)CuO\(_4\)\(^{27}\) and underdoped \(La_{2-x}Sr_xCuO_4\)\(^{35}\) and zinc-doped YBCO \([YBa_2(Cu_{1-x}Zn_x)O_8]\)\(^{36}\).

The two highest-frequency absorption peaks at around 30 000 – 40 000 cm\(^{-1}\) (4–5 eV) are typical for the cuprates and represent transitions into a band formed mostly by oxygen \(p\) orbitals (see, e.g., Ref.\(^{14}\)). The band slightly below 20 000 cm\(^{-1}\) (e.g., around 1.5 – 2 eV) is very similar to the upper Hubbard band. Such absorption bands have been observed, for example, in optical-conductivity studies of insulating undoped Pr\(_2\)CuO\(_4\)\(^{22}\) and Nd\(_2\)CuO\(_4\)\(^{22}\).

It is important to realize that the presence of such a band does not necessarily require a charge-transfer gap. Moreover, LDA + DMFT calculations\(^{44}\) demonstrated that such a band may perfectly coexist with a quasiparticle absorption peak (e.g., with a metallic state) in the case of undoped Nd\(_2\)CuO\(_4\) with a perfect \(T'\) structure.

C. Spectral weight

We can trace the temperature evolution of each term in Eq.\(^{2}\). The mid-infrared band is almost invariant with temperature, while the shape of the lower-frequency spectrum (consisting of the Drude contribution and the FIR band) changes. While at temperatures below 150 K the peak in \(\sigma_1\) is centered at \(\omega = 0\) (the Drude term...
dominates), it shifts to finite frequencies at $T > 150$ K. This shift is an indication of the breakdown of the simple Drude-metal picture. It suggests a continuous change in the charge transport from the low-temperature coherent (Drude) to high-temperature incoherent regimes.\(^\text{36}\)

In Fig. 6 we plot the spectral weight (SW) of the Drude term (panel a), the FIR band (panel b), the sum of the two (panel c), and, for completeness, the MIR band (panel d). (The spectral weight of each term in Eq. (A2) is just the squared plasma frequency of the term.) As one can see from Fig. 6 the spectral weights plotted in panels (c) and (d) are temperature independent, only the Drude and the FIR-band spectral weights depend on temperature. It is obvious that the spectral weight of the FIR band grows at the expense of the Drude component with increasing temperature. We suggest that this spectral weight transfer between the Drude and the FIR band is related to the change in the transport properties, namely to the change from the quadratic to the linear temperature dependence of $\rho(T)$ which happens at comparable temperatures (Fig. 1).

A qualitative picture of the spectral weight redistribution with temperature in PCO can be studied by means of the total spectral weight:

$$SW(\omega_c) = 8 \int_0^{\omega_c} d\omega \sigma_1(\omega).$$  \hspace{1cm} (3)

It is plotted as a function of the cutoff frequency $\omega_c$ in the inset on the right-hand panel of Fig. 5. At low frequencies $\omega_c$, $SW(\omega_c)$ increases with decreasing temperature, and it increases with $\omega_c$ for all temperatures, finally developing an upturn around 10000 – 15000 cm\(^{-1}\). This upturn is due to interband transitions. Up to 15000 cm\(^{-1}\), the spectral weight shows a temperature dependence. Only at higher frequencies do the spectral-weight curves merge, implying that above 15000 cm\(^{-1}\) (~1.9 eV) the spectral weight is conserved as temperature changes. In other correlated-electron materials, the spectral weight is known to be conserved also only at frequency scales of a few eV.\(^\text{4,39}\) Thus, our results indicate the presence of electron correlations in PCO.

In order to estimate the spectral weight and the plasma frequency of the itinerant charge carriers only, we set $\omega_c/2\pi = 9400$ cm\(^{-1}\), thus cutting off the contribution from the interband transitions. This gives a plasma frequency of 17700 cm\(^{-1}\) (2.19 eV), a value comparable to those for other high-$T_c$ cuprates.\(^\text{27,40,41}\) Using the relation between the charge-carrier density $n$ and the plasma frequency ($\omega_p^2 = 4\pi ne^2/m$), $n$ is estimated to give $\sim 3.53 \times 10^{21}$ cm\(^{-3}\) assuming $m$ to be equal to the free-electron mass $m_0$.

**D. Extended-Drude analysis**

To get further insight into the physics behind the optical response of Pr\(_2\)CuO\(_4\), we analyze the optical conductivity data in terms of the extended (or generalized) Drude model which is widely used for analysis of the optical properties of correlated electron systems.\(^\text{42,43}\) The complex conductivity in this model is given by

$$\sigma(\omega) = \frac{1}{4\pi} \frac{\omega_p^2}{\Gamma(\omega) - i\omega [1 + \lambda(\omega)]},$$  \hspace{1cm} (4)

where $[1 + \lambda(\omega)] = m^*(\omega)/m$ and $\tau_{op}^{-1}(\omega) = \Gamma(\omega)$ are the frequency-dependent mass renormalization factor and the optical scattering rate, respectively. Inverting Eq. (4) gives

$$1 + \lambda(\omega) = \frac{\omega_p^2}{4\pi \omega [\sigma(\omega)]^2}; \quad \Gamma(\omega) = \frac{\omega_p^2 \sigma_1(\omega)}{4\pi [\sigma(\omega)]^2}.$$  \hspace{1cm} (5)

The frequency-dependent optical scattering rate, obtained on the basis of Eq. (4) with $\omega_p = 2.19$ eV, is displayed in Fig. 7 as a function of frequency for various temperatures listed in the legend. At $T < 150$ K, the general trend in $\tau_{op}^{-1}(\omega)$ is to increase with frequency, but this increase is nonmonotonic. This is due to phonons and the localization mode discussed above. This mode reveals itself as a bump at around 230 cm\(^{-1}\) (~28 meV) in the optical scattering rate. At $T > 150$ K, the scattering rate increases rapidly as $\omega \to 0$. This is because at high temperatures the localization mode dominates the Drude contribution as it was discussed in relation to the $\sigma(\omega)$ spectra.

**E. Eliashberg analysis and electron-boson spectral density**

The optical scattering rate is according to App. B closely related to the electron-exchange boson interaction spectral density $I^2 \chi(\omega)$ [Eq. (B1)] which is at the core of
normal and superconducting state Eliashberg theory. This theory can be applied to calculate various normal and superconducting state properties and, consequently, it is of quite some interest to gain knowledge on \( I^2(\omega) \) by inverting \( \tau_{\text{op}}^{-1}(\omega) \). This will allow a more detailed analysis of our experimental results.

It was also demonstrated by Schachinger et al.\textsuperscript{44} that any nonzero contribution to \( I^2(\omega) \) at some energy \( \omega \) will result in an increase of the optical scattering rate. Consequently, the bump observed in the optical scattering rate of PCO (Fig. 7) at around 230 cm\(^{-1} \) (\( \sim 28 \) meV) cannot be caused by electron-exchange boson interaction and is, therefore, not part of the conducting-electron background. Nevertheless, we concentrate on the normal-state \( T = 30 \) K data and perform a straightforward inversion using the maximum-entropy procedure outlined in App. 13 by inverting Eq. 12 together with the kernel Eq. 15. As the temperature and frequency independent impurity scattering rate \( \tau_{\text{imp}}^{-1} = 2\pi t^+ \) is not known, this is an iterative process which is performed by slowly increasing \( t^+ \) until a smooth function \( I^2(\omega) \) with no pronounced spikes in the immediate vicinity of \( \omega = 0 \) has been found. This resulted in \( \tau_{\text{imp}}^{-1} \sim 100 \) meV (\( t^+ = 15 \) meV) which is quite substantial but in good agreement with what has been reported for the system PCCO.\textsuperscript{45} Furthermore, we restricted the frequency range of the inversion to \( \omega \in [0,300] \) meV because between 100 \( \leq \omega \leq 300 \) meV, \( \tau_{\text{op}}^{-1}(\omega) \) develops only a moderate increase with energy.

It has to be pointed out, though, that Eq. 12 is only approximate. Therefore, we use the spectrum \( I^2(\omega) \) which resulted from the inversion process to calculate the quasiparticle self-energy using the full normal-state infinite-bandwidth Eliashberg equations. The complex infrared conductivity \( \sigma(\omega,T) \) is then calculated using a Kubo formula\textsuperscript{43} and the resulting optical scattering rate is calculated from Eq. 1. A comparison of this result with the data requires some adaptation of the original \( I^2(\omega) \) spectrum in order to achieve the best possible agreement with the data. This final spectrum is presented in the inset of Fig. 7. It shows a double-peak structure which is followed by a deep valley and a hump at higher frequencies. The low-energy peak is at \( \sim 11 \) meV and the high-energy peak can be found at \( \sim 50 \) meV. Similar double-peak spectra have been reported for PCCO by Schachinger et al.\textsuperscript{46} and for La\(_{0.85}Sr_{0.15}CuO\(_4\) (a hole-doped cuprate) by Hwang et al.\textsuperscript{47} both with a less pronounced low-energy peak. It is most likely that the bump around \( \sim 28 \) meV in the PCO \( \tau_{\text{op}}^{-1}(\omega) \) is responsible for this overpronouncement of the low-energy peak in the PCO \( I^2(\omega) \) spectrum.

We found that the mass renormalization factor, which can be calculated as the first inverse moment of \( I^2(\omega) \), is \( \lambda = 4.16 \).

A comparison of theoretical and experimental \( \tau_{\text{op}}^{-1}(\omega) \) data for \( T = 30 \) K is presented in Fig. 7(b). The solid (red) curve represents the data while the dashed (black) curve presents the result of our theoretical calculations on the basis of the \( I^2(\omega) \) spectrum shown in the inset. (Of course, good agreement between theory and data cannot be expected in the energy region around the bump at \( \sim 28 \) meV.)

F. Penetration depth and Superfluid density

The temperature dependence of the penetration depth of PCO was obtained experimentally by means of phase-sensitive millimeter-wave measurements.\textsuperscript{25} Using the Fresnel optical formulas for the complex transmission coefficient, the in-plane complex conductivity of the film was calculated directly from the measured transmission coefficient and phase shift. The penetration depth was then calculated from \( \sigma_2 \) by using \( \lambda_L = c/(4\pi\omega\sigma_2)^{1/2} \), where \( c \) is the vacuum speed of light and \( \omega \) is the frequency of the incoming radiation. We found \( \lambda_L(T \to 0) \approx 1.6 \pm 0.1 \) \( \mu m \).

Figure 8(a) presents the normalized superfluid density \( N_s(T) = n_s(T)/n_s(0) = \lambda_L^2(0)/\lambda_L^2(T) \) measured at 7 cm\(^{-1} \) [solid (red) circles] and 8.3 cm\(^{-1} \) [solid (blue) triangles] vs temperature. We added curves for \( N_s(T) = 1 - (T/T_c)^2 \) [thin dashed (purple) curve] and \( N_s(T) = 1 - (T/T_c)^4 \) [thin dashed-dotted (olive) curve] for comparison. They are supposed to mimic the temperature dependence of \( N_s(T) \) for nodal (d-wave) and fully gapped (s-wave) superconductivity, respectively. Obviously, the former curve describes the data reasonably well, whereas the fully gapped behavior can certainly be ruled out.

More sensitive is the low-temperature variation of the penetration depth \( [\lambda_L(T) - \lambda_L(0)]/\lambda_L(0) \) as a function of...
FIG. 8: (Color online). Panel (a): Superfluid density, $N_s(T)/\lambda_{c}(T)$, as a function of temperature. Panel (b): Low-temperature variation of the normalized London penetration depth, $|\lambda_L(T) - \lambda_L(0)|/\lambda_L(0)$, as a function of temperature squared. Data derived from the millimeter-wave conductivity measurements at 7 cm$^{-1}$ and 8.3 cm$^{-1}$ are presented by solid (red) circles and solid (blue) triangles, respectively. Panel (a) contains for comparison the temperature dependence $N_s(T) = 1 - (T/T_c)^{\gamma}$ [thin dashed (purple) line] which is expected for a nodal superconductor and $N_s(T) = 1 - (T/T_c)^{\gamma}$ [thin dashed-dotted (olive) line] for a fully gapped superconductor. In panel (b) a quadratic power law of the reduced penetration depth is indicated by a thick solid (black) line. Inset: A universal relation between the zero-temperature superfluid density, $N_{s0}$, and the product of normal-state dc conductivity and $T_c$ as found in Ref. 52 [straight solid (blue) line], reported “error bars” of this relation [straight dashed (blue) lines], and the data obtained for three PCO films investigated in this study and in Ref. 51 [bold (red) circles]. The error bars are shown for the least accurate data.

the square of the temperature. The data are presented in Fig. 8(b) using the same symbols as in Fig. 8(a). For a nodal superconductor a quadratic power law [thin solid (black) line] is to be expected and the data are in agreement with this power law at lowest $T$.

In returning to the absolute value of the zero-temperature penetration depth $|\lambda_L(T \to 0)| \approx 1.5 \pm 0.1$ km we conclude that the density of the superfluid condensate is very low in PCO as compared to typical values for optimally doped cuprates, where the penetration depth is smaller by a factor of 5 to 10. For example, in optimally doped PCCO $\lambda_L = 330$ nm. This points toward a doping-related nature of superconductivity in our PCO samples, as large values of $\lambda_L(0)$ are typical for either underdoped or overdoped regimes. However, the value of $\lambda_L(0)$ found here is so big, that within this picture our PCO sample must be far off optimal doping. This is rather unlikely, because the critical temperature of our film is definitely too high for a heavily underdoped or overdoped sample. It is also important to note that a possible degradation of $T_c$ while the optical measurements have been performed can be excluded because the dc resistivity measured after the completion of all optical measurements does not differ from the resistivity measured on the fresh film (see bottom panel of Fig. 8).

Furthermore, we would like to note that our second PCO sample had a $\lambda_L(0) = 1.55 \pm 0.25$ µm and a very similar value of $T_c$. Another PCO film prepared by a different method (metal organic decomposition vs MBE for the current films) was reported by some of us to have $\lambda_L(0) = 1.35 \pm 0.25$ µm and $T_c = 27.5$ K.

It is revealing to note that all these samples do not fit a supposedly universal relation reported by Homes et al. which connects the superfluid density [or $\lambda_L(0)$] of cuprates to the product of $T_c$ and normal-state conductivity $\sigma_{dc}$. This relation was obtained from an analysis of experimental data on doped samples, e.g. on doped charge-transfer insulators, and works for all doping levels. According to Zaanen the existence of this universal relation reflects the fact that the normal state of the doped cuprates is extremely viscous (dissipative).

All the PCO samples studied here and in Ref. 51 are far off this universal relation (see the inset in Fig. 8). It is tempting to explain this fact as a sign of a possible departure from the charge-transfer-insulator picture in PCO while Homes’ relation reflects the physics behind the doped-insulator picture, it does not necessarily work any longer whenever this picture loses its validity in cuprates.

Nevertheless, if our samples are indeed doped, it seems to be reasonable to assume that they must be underdoped rather than overdoped. This assumption is based on the method used to prepare the samples (reduction of oxygen content), on the high value of $T_c$, and on the low superfluid density.

G. Absence of a pseudogap feature

A well-known characteristic feature of the underdoped cuprates is the occurrence of a pseudogap, i.e. a partial normal-state gap in the electronic density of states. Such a pseudogap has been observed in underdoped cuprates by many experimental methods. In optical experiments, the occurrence of a pseudogap below a characteristic temperature can manifest itself in different ways. In hole-doped cuprates, the pseudogap is seen as a suppression of the low-frequency scattering rate. In electron-doped cuprates a suppression of the MIR reflectivity is observed that corresponds to a reduced real-part MIR optical conductivity and to a nonmonotonic behavior of the so-called restricted spectral weight, $RSW(\omega_L, \omega_H, T)$. Here, RSW is defined as

$$RSW(\omega_L, \omega_H, T) = 8 \int_{\omega_L}^{\omega_H} d\omega \sigma(\omega, T),$$

where $[\omega_L, \omega_H]$ is the restricted frequency range of interest.

It follows from the reflectivity and conductivity data presented in Figs. 2 and 4 that such a normal-state gap
is not evident in the current system. The absence of the normal-state pseudogap is further confirmed by the results presented in Fig. 9. In this figure, the restricted spectral weight RSW(ωL, ωH, T) normalized to the spectral weight at T = 300 K is plotted as a function of temperature for four frequency ranges [ωL, ωH] as quoted in the legend. It is evident that the normalized restricted spectral weight displays in all four cases a monotonic temperature dependence. This rules out the existence of a normal-state pseudogap. In addition, the optical scattering rate (see Fig. 10) shows (apart from the low-frequency features due to localization, phonon modes, and MIR bands) no temperature-dependent suppression, that might (similarly to the hole-doped cuprates) indicate a pseudo-gap-like feature.

Thus, we conclude that a pseudogap is absent in the PCO films, in contrast to most underdoped high-Tc cuprates. In our view, this difference can be related to the absence of an antiferromagnetic phase in PCO. In electron-doped cuprates, the magnetic order induces the pseudogap. With doping, the Neel temperature decreases monotonically leading to a complete suppression of antiferromagnetic order at higher doping accompanied by the disappearance of the pseudogap. The absence of a pseudogap feature in PCO supports ideas, expressed in Refs. 10, 12, about a strong suppression or even absence of an antiferromagnetic insulating phase in electron-doped cuprates, if the T' structure (i.e., no apical oxygen) can be managed to survive down to very low or even zero doping levels.

V. CONCLUSIONS

In our broadband investigation of the optical response of thin PCO films, we unveiled a low-frequency (FIR) collective charge excitation attributed to localization effects. As a function of temperature, the optical spectral weight is redistributed between this mode and the zero-frequency-centered Drude peak: the weight of the FIR mode grows with temperature at the expense of the Drude-peak weight. Such a behavior has been reported in underdoped cuprates and is typical for bad metals.

We found that the optical spectral weight remains temperature-dependent up to 1.9 eV, which indicates strong electron correlations in PCO.

We calculated the electron-boson spectral density and found the mass renormalization factor, λ = 4.16 at 30 K.

In the millimeter-wave data, we directly observed the formation of the superconducting condensate at T < Tc. We obtained that the temperature dependence of the London penetration depth at low temperatures follows a quadratic power law. This indicates d-wave symmetry which is typical for the cuprates.

Neither the experimental optical data nor their analysis reveal any indication of normal-state gap-like features which could be attributed to the existence of a normal-state pseudogap. This observation is in line with a breakdown of the charge-transfer-insulator picture in PCO.

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Appendix A: Thin Films Optical Conductivity

In this work we measured a thin PCO film on top of a DyScO3 substrate. Harbecke studied the optical response of multilayer systems in the most general case, including so-called “coherent” and “incoherent” light propagation.

Coherent propagation is expected when the optical thickness of a medium is comparable or smaller than the wavelength of light. Our measurements span wavelengths from the far-infrared to the ultraviolet. The film thickness is of the order of a few hundred angstroms. We can safely assume coherent propagation throughout the whole measured range. Dealing with the substrate is trickier. Our substrate is 0.5 mm thick. In the far-infrared, light propagates coherently but coherence is gradually lost when moving into the mid-infrared and shorter wavelengths.

To solve this problem, we worked with an unpolished back surface of the substrate. The roughness of our back surface is of the order of 10–50 μm, strongly diffusing light of shorter wavelengths. For all that matters, light reflected from the back surface will never reach the detector for this range. A problem might still arise in the far-infrared. However, we have a highly metallic, hence absorbing, film. The contribution from the back surface of the substrate becomes negligible as it implies going twice through the film and being partially diffused by the back surface.
In practice, the effective geometry that best describes our system is a thin film (thickness $d$ and complex refraction index $n_f$), where light propagates coherently, sitting on top of a half-infinite substrate having a complex refraction index $n_s$. The near normal incidence reflectivity of such a system is

$$R = \left| r_0 + \frac{n_f^2 r_f \phi_f^2}{1 + r_f r_0 \phi_f^2} \right|^2,$$

(\text{A1})

where $r_0 = (1 - n_f)/(a + n_f)$, $t_0 = 2/(1 + n_f)$, $r_f = (n_f - n_s)/(n_f + n_s)$, and $\phi_f = \exp(2\pi i n_f d/\lambda)$, $\lambda$ being the wavelength of light.

In principle, one can perform Kramers-Kronig analysis on the measured reflectivity and numerically invert Eq. (A1) to obtain $n_f$, assuming that we measured the substrate ($n_s$) independently. In practice this procedure is strongly dependent on initial trial values and does not converge with reasonable accuracy. An alternative approach is to model the bulk dielectric function ($\varepsilon = n_f^2 = 4\pi \sigma/\omega$) of the film material. We enter this model into Eq. (A1) and utilize a least squares fitting to refine its parameters.

The most straightforward modeling for the dielectric function is the Drude-Lorentz formalism:

$$\varepsilon_{DL} = \varepsilon_{\infty} - \frac{\omega_{pD}^2}{\omega^2 + i\sigma \tau^{-1}} + \sum_k \frac{\omega_{pk}^2}{\omega_{nk}^2 - \omega^2 - i \gamma_k \omega},$$

(\text{A2})

In Eq. (A2) $\varepsilon_{\infty}$ is the contribution from electronic transitions in the deep-UV. The second term corresponds to a free-carrier Drude response, characterized by a plasma frequency $\omega_{pD}$, and a frequency-independent scattering rate $\tau^{-1}$. The last term is a sum of Lorentz oscillators; each of them is characterized by a resonance frequency $\omega_{nk}$, a line width $\gamma_k$, and a plasma frequency $\omega_{pk}$. All these parameters may freely vary with temperature. This approach has been successfully utilized in several cuprate thin films and a detailed analysis can be found in Santander-Syro et al.\textsuperscript{64}

However, the use of Eq. (A2) is model dependent. We went a step further in our analysis by adapting the procedure proposed to extract optical functions from single crystal data by Kuzmenko.\textsuperscript{65} In his approach the reflectivity is roughly fitted by a Drude-Lorentz dielectric function. A second step is then taken by choosing a variational dielectric function $\varepsilon_V$. This function is added to $\varepsilon_{DL}$ and adjusted in order to describe the total reflectivity within data noise. In Kuzmenko’s paper, $\varepsilon_V$ is obtained by setting an arbitrary piece-wise imaginary part ($\varepsilon''$) of the dielectric function, and calculating the corresponding real part from Kramers-Kronig. This piecewise $\varepsilon''$ is obtained by simply setting an arbitrary value at each measured frequency. This value is modified in a least-squares fit so that the reflectivity is properly adjusted. Kuzmenko showed that this method produces optical functions with an accuracy equivalent to Kramers-Kronig.

In principle, Kuzmenko’s method can be straightforwardly adapted to the case of thin films if one replaces the equation for the normal-incidence reflectivity in bulk materials by Eq. (A2). However, because of the very large spectral range of our data setting an arbitrary value for $\varepsilon''$ at each point measured would be impractical. What we did, instead, was to add a very large number (of the order of 1000) of Lorentz oscillators distributed over the regions where the first fit by the Drude-Lorentz model does not describe properly the data. We fixed the frequency and width of these oscillators and allowed its intensity to vary. In particular, we modified the form shown in Eq. (A2) in order to allow for negative values of the numerator. These modified Lorentz oscillators should not be regarded as physical excitations of the systems. They represent a differential correction on the rough Drude-Lorentz dielectric function.

At the end of the procedure, we have a dielectric function

$$\varepsilon = \varepsilon_{DL} + \varepsilon_V,$$

(A3)

which is a model-independent description of the bulk optical properties of the thin-film material. As a bonus, the rough $\varepsilon_{DL}$ gives us an overall idea of the physical excitations in the system. However, the obtained dielectric function is model-independent and can be used even when the Drude-Lorentz approach is not expected to work, such as in the superconducting state. Furthermore, usage of the Drude term is not a must even in a metallic state. One could use a large number of Lorentz oscillators instead.

**Appendix B: Maximum entropy inversion**

The optical scattering rate $\tau_{op}^{-1}(\omega)$ is directly related to the electron-exchange boson interaction spectral density (Eliashberg function) $I^2\chi(\omega)$ via\textsuperscript{45,62}

$$\tau_{op}^{-1}(\omega,T) - \tau_{imp}^{-1} = \int_0^\infty d\Omega K(\omega,\Omega,T)I^2\chi(\omega).$$

(\text{B1})

The kernel $K(\omega,\Omega,T)$ is determined from theory, $\tau_{imp}^{-1}$ is an impurity scattering rate and $I^2\chi(\omega)$ can be calculated by deconvoluting (inverting) Eq. (B1). Shulga et al.\textsuperscript{53} derived for the normal state the kernel

$$K(\omega,\Omega,T) = \frac{\pi}{\omega} \left[ 2\omega \coth \left( \frac{\Omega}{2\pi} \right) - (\omega + \Omega) \coth \left( \frac{\omega + \Omega}{2T} \right) \right. \left. + (\omega - \Omega) \coth \left( \frac{\omega - \Omega}{2T} \right) \right].$$

(B2)

This kernel is a very good approximation to the exact result of Eliashberg theory. It is valid for any temperature $T$ and reduces to the kernel derived by Allen\textsuperscript{64} for $T = 0$.

The maximum-entropy method to invert Eq. (B1) to gain information on $I^2\chi(\omega)$ from experimental $\tau_{op}^{-1}(\omega,T)$
data was described in detail by Schachinger et al.\textsuperscript{12} This process is, of course, an ill-posed problem which does not necessarily allow for an unique solution.

Let us define
\[ \chi^2 = \sum_{i=1}^{N} \frac{(D_i - \tau_{iop}^{-1}(\omega_i))^2}{\varepsilon_i^2}, \]  
(B3)

where \( D_i \) are the experimental \( \tau_{iop}^{-1} \) data points at discrete energies \( \omega_i \) and \( \tau_{iop}(\omega) \) is calculated from Eq. (B1) and kernel Eq. (B2) and is to be regarded as a functional of \( I^2\chi(\omega) \). Finally, \( \varepsilon_i \) denotes the error bar on the data \( D_i \) and \( N \) is the number of data points. Furthermore, physics requires that the spectral density \( I^2\chi(\omega) \) is positive definite. To achieve this the maximum-entropy method minimizes the functional
\[ L = \chi^2 - aS, \]  
(B4)

with \( S \) the generalized Shannon-Jones entropy\textsuperscript{45,46}
\[ S = \int_0^\infty d\omega \left\{ I^2\chi(\omega) - m(\omega) - I^2\chi(\omega) \ln \frac{I^2\chi(\omega)}{m(\omega)} \right\}, \]  
(B5)

which gets maximized in the process. \( m(\omega) \) is the constraint function (default model) which reflects \textit{a priori} knowledge of \( I^2\chi(\omega) \). In Eq. (B4) \( a \) is a determinative parameter that controls how close the fitting should follow the data while not violating the physical constraints. In our inversion we set \( m(\omega) = m_0 \) for \( \omega_1 \leq \omega \leq \omega_N \) with \( m_0 \) some small constant indicating that we have no knowledge whatsoever about \( I^2\chi(\omega) \), thus establishing an \textit{unbiased} inversion of Eq. (B1). Finally we will make use of the historical maximum-entropy method which iterates \( a \) until the average \( \langle \chi^2 \rangle = N \) is achieved with acceptable accuracy.

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