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Publication Date
1970-04-01
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April 1970

AEC Contract No. W-7405-eng-48

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Phonon Contribution to the Far Infrared Absorptivity of Superconducting and Normal Lead

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ABSTRACT

Low temperature measurements of the absorptivity of superconducting and normal Pb in the phonon frequency region 15-200 cm$^{-1}$ show an onset of absorption when the photon energy is large enough to excite appreciable numbers of acoustic phonons. The onset begins around the transverse phonon frequency $\Omega \approx 35$ cm$^{-1}$ in the normal metal and around $\Omega + 2\Delta \approx 55$ cm$^{-1}$ in the superconductor. Most features of the data can be explained by a Golden Rule calculation of the phonon generation process proposed by Holstein.
We have measured the far infrared absorption in single crystals of pure lead over the frequency range from $15 \text{ cm}^{-1}$ to $200 \text{ cm}^{-1}$ in both the superconducting and normal states. These data show the first example of structure on the absorptivity of a metal due to the excitation of phonons. It is helpful to consider the analogy to the phonon sidebands commonly observed on electronic transitions in insulators. In our case the "electronic transition" is the excitation spectrum of the normal or superconducting metal, so an increase in absorption due to phonon creation is expected to start at characteristic phonon frequencies in the normal metal and at the phonon frequencies plus $2\Delta$ in the superconductor. The absorption spectra obtained increase dramatically at the expected frequencies, and structure is observed which can be correlated with peaks in the density of phonon states in Pb. We have thus observed the onset of the volume phonon generation process proposed by Holstein to account for the near infrared absorptivity of metals.

In our experiment radiation from a far-infrared Michelson interferometer was focused onto a single crystal sample of lead about $7\times7\times0.5 \text{ mm}$ in size mounted in a highly absorbing cavity. A doped germanium thermometer was cemented to the back of the sample to monitor its temperature. The sample was supported in the cavity by thin-walled stainless steel tubing which served the dual purposes of thermally isolating the sample and protecting the thermometer from directly absorbing any radiation reflected from the sample. The samples had a residual resistance ($R_{300}/R_T$) of 13000 at $4.2^\circ\text{K}$ and 85000 at $1.2^\circ\text{K}$. Their surfaces were carefully chemically polished so as to maximize the specular reflection of visible light. The experiments done at $1.2^\circ\text{K}$ on samples in the superconducting state were compared with similar
experiments on normal state lead done in a magnetic field of ~ 1200 Gauss applied parallel to the sample surface. Reference spectra were obtained by substituting a highly absorbing carbon resistance bolometer for the lead sample. Analysis of the data consisted of calculating the ratios of the lead spectra to the carbon bolometer spectrum and the ratio of the superconducting lead absorption spectrum to the normal absorption spectrum. The superconducting/normal absorption ratios are more useful because any spurious structure due to resonant cavity modes should cancel out of this ratio. The results of many experiments on several samples in two different cavities were averaged and standard deviation confidence levels were computed. No significant differences were observed in the spectra obtained for (100) and (111) sample orientations. This is to be expected since the frequencies are generally too high for the anomalous skin effect to retain appreciable Fermi surface selectivity.

Figure 1 shows the absorption spectra for superconducting and normal samples normalized to the carbon bolometer. The solid line is the frequency-dependent (surface) absorption arising from the ordinary skin effect theory. This was calculated from Dingle's tables using the relaxation time derived from the resistance ratio measurement and Chambers' value of $\sigma/\lambda$. The shape of the theoretical curve is not very sensitive to errors in the chosen parameters. The absolute accuracy of the measurements is poor so the scale factor is established by fitting the curves in the region from 20-40 cm$^{-1}$ where the theory is expected to be valid.

The spectrum for the superconducting sample shows a well-defined
energy gap absorption edge at 22 cm$^{-1}$ with very little signal at lower frequencies; this indicates that at least 95% of the signal is due to absorption in the lead sample. In addition, the absorption increases sharply at about 55 cm$^{-1}$. A similar but less sharp increase begins at about 35 cm$^{-1}$ in the normal sample. Although the onset of the additional volume absorption is slow in the normal state, it must be invoked to explain the fact that the spectrum becomes concave upward. The skin effect theory predicts a curve which is concave downward throughout this frequency region. These curves show that the Holstein absorption is comparable to the skin effect absorption for frequencies above the phonon range.

The details of the onset of absorption are seen more clearly in Fig. 2, where the ratio of superconducting to normal absorption is plotted. The ratio rises rapidly from the gap at 2$\Delta$=22 cm$^{-1}$ to a maximum at 35 cm$^{-1}$. The decrease from 35-55 cm$^{-1}$ is due to the absorption onset in the normal spectrum. The ratio then rises in two steps, centered at $\sim$ 65 and $\sim$ 95 cm$^{-1}$, which are due to the onset of absorption in the superconducting spectrum.

The accuracy of the data in Fig. 1 decreases toward each end of the measured frequency range. Error bars are shown which indicate that the details of the structure around 150 cm$^{-1}$ are probably not significant. Many errors, both random and systematic, are expected to disappear from the ratio shown in Fig. 2. The data shown there are probably accurate to $\pm$ .02. Thus, the fine structure near 30 cm$^{-1}$ and beyond 100 cm$^{-1}$ is probably not significant, while some of the sharp features near 85 cm$^{-1}$ may be.

As an independent check of our absorption measurements, we measured the power reaching a bolometer after many reflections in a
cavity whose walls were slabs of single crystal Pb, a cavity used previously to study energy gap absorption edges in superconductors. The ratio of superconducting to normal absorptivity derived from this experiment was in good agreement with our direct absorption data.

The Holstein process involves the scattering of a conduction electron which simultaneously absorbs a photon and emits a phonon. A rigorous treatment of this process requires a knowledge of the Fermi surface and phonon spectrum of Pb, as well as detailed consideration of the superconducting state. A phenomenological approach suggested by Falicov\(^9\) can be used, however, to understand the main features of the data. This approach assumes the conduction electrons to be a free electron gas at T=0 and includes energy conservation but not wave-vector conservation selection rules. The initial state is a photon of energy \(\omega\); the final state is an electron of energy \(\varepsilon_1\), a hole of energy \(\varepsilon_2\), and a phonon of energy \(\Omega\). The density of final states is assumed to be the convolution of the independent densities of states of the electron, hole, and phonon. The matrix element is assumed constant except for a factor \(\omega^{-1/2}\) which arises from the initial photon state, and a possible dependence of the electron-phonon interaction on the phonon frequency. The volume absorptivity is thus proportional to

\[
\mathcal{A}_v = \frac{1}{\omega} \int N_e(\varepsilon_1) N_h(\varepsilon_2) \alpha^2(\Omega) F(\Omega) \delta(\varepsilon_1 + \varepsilon_2 + \Omega - \omega) \, d\varepsilon_1 \, d\varepsilon_2 \, d\Omega
\] (1)

where \(N_e\) and \(N_h\) are the electron and hole densities of states, and \(\alpha^2(\Omega) F(\Omega)\) is the product of the square of the electron-phonon matrix element and the phonon density of states. Substitution of a constant
for the normal electron density of states and a BCS density of states in the superconductor gives the expressions:

\[ a_{vN}(\omega) = \frac{N^2}{\omega} \int_0^{\omega} (\omega - \Omega) a^2 F(\Omega) d\Omega \quad (2) \]

\[ a_{vS}(\omega) = \frac{N^2}{\omega} \int_0^{\omega} (\omega - 2\Delta) \left[ (\omega - \Omega) E(a) - \frac{2\Delta^2}{(\omega - \Omega) K(a)} \right] d\Omega \quad (3) \]

where \( E(a) \) and \( K(a) \) are complete elliptic integrals with \( a^2 = 1 - \frac{4\Delta^2}{(\omega - \Omega)^2} \).

These integrals were calculated using values of \( a^2(\Omega) F(\Omega) \) obtained from superconducting tunneling. A composite spectrum was then calculated by adding the superconducting or normal theoretical Holstein volume absorptivity \( a_v \) to the theoretical anomalous skin effect absorptivity \( a_s \). The ratio of these two absorptivities was estimated by Holstein to be \( a_v/a_s = 16 \Theta_D \delta_f / 15 T \tau_f \) in the limit \( \hbar \omega \gg k \Theta_D \) where both \( a_v \) and \( a_s \) are independent of frequency. Here \( \delta_f \) is the high frequency skin depth \( c/\omega_p \), and \( T \) is the high temperature, large compared to \( \Theta_D \), at which the phonon limited relaxation time \( \tau \) is evaluated. The composite absorption in this high frequency limit has been used successfully to fit the near infrared absorptivity in copper and silver at low temperatures. Using the experimental value of \( \sigma/\lambda \) and the London penetration depth \( c/\omega_p \) for Pb we estimate \( a_v/a_s \approx 1.9 \). A reasonable fit to the data in Figs. 1 and 2, however, requires the smaller value \( a_v/a_s \approx 1 \).

The theoretical ratios of the superconducting and normal spectra are compared with the data in Fig. 2. The two curves shown for the theoretical ratio differ by the way in which the conventional skin
effect contribution to the absorption was estimated. In B the correct normal state skin effect from the Dingle tabulation of the theory was used for both the normal and superconducting states. Energy gap effects on the superconducting skin effect absorption are ignored. The shape of curve B clearly accounts for many of the major features of the data. The computed curve drops smoothly above 35 cm$^{-1}$ due to the onset of absorption in the normal state obtained from Eq. (2). It then rises in two steps due to the onset of absorption in the superconducting state. The sharply peaked BCS density of electronic states at the energy gap weights the phonon density of states for $\Omega$ near $\omega - 2\Delta$. Thus the peaks in $\sigma^2 F(\omega - 2\Delta)$ shown in curve D contribute step increases to Eq. (3) and thus to $A_S/A_N$.

Although qualitatively correct, the structure predicted from our simple model deviates from the measurements in that the latter are systematically broadened and shifted to higher frequencies. For example, the theoretical absorption steps centered at $-59$ and $-88$ cm$^{-1}$ appear in the measurements at $-65$ and $-95$ cm$^{-1}$ respectively. This is an understandable consequence of wave vector selection rules neglected in the theory. Eq. (1) implicitly assumes that any electron momentum is available, a good but not perfect approximation for Pb.

In curve C the skin effect absorption was computed in the extreme anomalous limit from values of $\sigma_1(\omega)$ and $\sigma_2(\omega)$ calculated from strong coupling superconductivity theory for Pb. The extreme anomalous limit is not accurate for our experimental conditions; it is known, for example, to underestimate the steepness of the onset of $A_S$ above the gap. The more accurate wave vector dependent calculations of
$A_S/A_N$ by Shaw and Swihart\textsuperscript{13} are in excellent agreement with the experimental gap edge, but do not extend to the phonon frequency region. Despite this inaccuracy, curve C illustrates two important points. The complex conductivities calculated from strong coupling superconductivity theory show structure due to virtual phonon effects. This structure appears as small (2-4\%) peaks in the skin effect absorption centered above the peaks in $\alpha^2 F(\omega-2\Delta)$ (curve D). These peaks are small compared with the steps due to real phonon generation and are not identifiable in curve C or in the data. A second important effect\textsuperscript{13} (a dispersion in the gap function) causes a broad dip in curve C between 100 and 300 cm\textsuperscript{-1}. The fall in the data beyond 120 cm\textsuperscript{-1} may thus be due to strong coupling effects. Preliminary measurements at higher frequencies show that $A_S/A_N$ continues to fall until 210 cm\textsuperscript{-1} and then remains constant to 350 cm\textsuperscript{-1}. This shape, but not the magnitude, is in agreement with our approximate theory.

ACKNOWLEDGEMENTS

The authors are greatly indebted to Professor L. M. Falicov for suggesting the simple model used to estimate the theoretical Holstein absorption and to Professor J. C. Swihart for providing values of $\sigma_1$ and $\sigma_2$ used in calculating the skin effect absorption. Thanks are also due to Professors M. L. Cohen and D. Scalapino, and Dr. P. Allen for helpful discussions. Our renewed interest in this problem was stimulated by discussions with Dr. H. Scher.

This work was done under the auspices of the U.S. Atomic Energy Commission.
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6. Our data are in good agreement with the more recent measurements in the gap region. Unfortunately, previous work on bulk samples was not, in general, carried out to high enough frequencies (or with adequate accuracy) to clearly show the Holstein absorption.


12. Holstein's estimate can be written \( \frac{\alpha_v}{\alpha_s} = 16 \Theta_D \lambda_L / 15 \Omega \). We used Chambers' value\(^5\) of \( \alpha/\lambda = 9.4 \times 10^{10} \Omega^{-1} \text{ cm}^{-2} \), \( \lambda_L = 305 \text{ Å} \) from R.F. Gasparovic and W. L. McLean (to be published), and handbook values of \( \sigma T \) and \( \Theta_D \).
FIGURE CAPTIONS

Fig. 1. Measured frequency dependent absorptivity in superconducting and normal lead compared with the prediction of normal state anomalous skin effect theory. The limits of the error bars are plus and minus one standard deviation (75 percent confidence).

Fig. 2. Measured ratio (curve A) of the absorptivity in superconducting/normal lead compared with B) a theoretical estimate of $A_S/A_N$ which ignores the effect of the energy gap on the superconducting surface absorption, but treats the anomalous skin effect correctly, C) a second estimate of $A_S/A_N$ with the surface absorption taken from the extreme anomalous limit of strong-coupling superconductivity theory, and D) $\alpha^2(\omega-2\Delta)F(\omega-2\Delta)$ the square of the electron-phonon coupling constant times the phonon density of states shifted in frequency by the amount $2\Delta$. Curves B and C are plotted on the same scale as curve A, but have been shifted vertically for clarity.
Fig. 1
Fig. 2
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