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Terahertz conductivity at the Verwey transition in magnetite

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The complex conductivity at the (Verwey) metal-insulator transition in Fe₃O₄ has been investigated at terahertz and infrared frequencies. In the insulating state, both the dynamic conductivity and the dielectric constant reveal a power-law frequency dependence, the characteristic feature of hopping conduction of localized charge carriers. The hopping process is limited to low frequencies only, and a cutoff frequency νₗ ≈ 8 meV must be introduced for a self-consistent description. On heating through the Verwey transition the low-frequency dielectric constant abruptly decreases and becomes negative. Together with the conductivity spectra this indicates the formation of a narrow Drude peak with a characteristic scattering rate of about 5 meV containing only a small fraction of the available charge carriers. The spectra can be explained assuming the transformation of the spectral weight from the hopping process to the free-carrier conductivity. These results support an interpretation of Verwey transition in magnetite as an insulator-semiconductor transition with structure-induced changes in activation energy.

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I. INTRODUCTION

Magnetite (Fe₃O₄) is probably the oldest known magnetic material that can be found in natural form. This material is of considerable importance because of various applications in magnetic recording and for high-frequency electronic devices. Physical properties of magnetite have attracted much attention after the discovery of the first-order metal-insulator transition at Tᵥ ≈ 120 K. On cooling through the transition temperature, the dc conductivity drops by two orders of magnitude. A realistic model to explain the mechanism of this transition has been suggested by Verwey, assuming a charge order-disorder transition with alternating valence (Fe²⁺/Fe³⁺) of the octahedrally coordinated iron ions. The metal-to-insulator transition in magnetite has been termed the Verwey transition since then. In spite of a large number of experimental and theoretical efforts, the mechanism governing the conduction and magnetic properties in magnetite is still under debate. According to the results of recent x-ray resonant scattering, even the charge ordering below Tᵥ has been questioned, and the concept of an itinerant magnet was considered instead. Also, the importance of the orbital degrees of freedom has recently been highlighted.

Magnetite crystallizes in a cubic high-temperature structure and exhibits a monoclinic distortion below the Verwey transition. In spite of enormous progress in resolving the low-temperature monoclinic structure of magnetite, full details could not be completely resolved so far. The monoclinic phase is insulating and is characterized by a gap of the order of 100 meV in the excitation spectrum, which is seen in photoemission data, in thermoelectric properties, and by optical spectroscopy. The charge transport in magnetite is usually explained within a polaronic picture. However, there is no general agreement about the energy of the polaronic absorption in the conductivity spectra. Recent photoemission results, obtained in the same magnetite crystals as in the present paper, have been self-consistently explained using a small-polaron model. A similar concept has been applied recently to explain the dynamic conductivity at the metal-insulator transition in quasi-one-dimensional β-Na0.33V2O5.

At low temperatures the conduction in magnetite takes place via hopping between localized states, which agrees well with both activated dc resistivity and with an observed characteristic power law σ ∝ ν° (Ref. 21) of the ac conductivity. This universal power law has been observed in various materials with internal disorder and is a clear fingerprint of hopping of charge carriers. The situation gets more complicated in the metallic state. In spite of two orders-of-magnitude increase in resistance above Tᵥ, the absolute value of the dc conductivity [σdc(T ≥ Tᵥ) ~ 50 Ω⁻¹ cm⁻¹] is still much smaller than the Ioffe-Regel minimum metallic conductivity σmin ~ e²/3ℏa ~ 3000 Ω⁻¹ cm⁻¹, even including the Mott correction in the case of the disorder σmin ~ 0.03e²/ℏa ~ 300 Ω⁻¹ cm⁻¹ (here a ~ 3 Å is the interatomic distance). The high-temperature conductivity of magnetite is thus typical for a semiconductor with a high mobility of thermally excited charge carriers rather than for a metal. Reviews of the recent and past developments in magnetite can be found in Refs. 2, 3, and 28–30.

In this paper we investigate terahertz (THz) and infrared conductivity of magnetite on both sides of the metal-to-insulator transition. The unexpected result of these experiments is the abrupt change of the dielectric constant at the...
transition temperature yielding even negative values above \(T_v\). Together with the conductivity spectra, these results indicate the formation of a narrow band of quasi-free carriers that contains only a small part of the total spectral weight.

II. EXPERIMENTAL DETAILS

Synthetic single crystals of magnetite were prepared from \(\alpha-\text{Fe}_2\text{O}_3\) using a floating-zone method with radiation heating.\(^3\) Two single crystals were used for the measurements, which revealed slightly different transition temperatures, \(T_v\approx 123\pm1\) K and \(116\pm1\) K, respectively. The transition temperature in magnetite can be taken as a criterion for impurity concentration and oxygen stoichiometry,\(^2,32\) which, therefore, differ significantly for the two crystals investigated. Both samples revealed qualitatively similar electrodynamic properties, revealing only differences in absolute values. Therefore, in the following, mainly the results on a sample with \(T_v\approx 123\) K will be shown. Plane-parallel samples of different thicknesses between 0.5 and 0.05 mm have been prepared from the original boules. To ensure mechanical stability, the thinnest samples were glued onto a MgO substrate of \(\approx0.5\) mm thickness. The availability of samples with different thicknesses is essential for transmittance experiments due to the strongly temperature-dependent absorption in magnetite.

The dynamic experiments for frequencies \(4\) cm\(^{-1}\leq\nu<40\) cm\(^{-1}\) were carried out in a Mach-Zehnder interferometer arrangement,\(^33\) which allows both: the measurements of the transmittance and the phase shift of a plane-parallel sample. The complex transmission coefficient has been analyzed using the Fresnel optical formulas for transmittance \(T=|t|^2\) of a plane-parallel sample:

\[
t = \frac{1 - r^2}{1 - 2r \cos \phi + r^2},
\]

where \(r=(\sqrt{e^*}-1)/(\sqrt{e^*}+1)\) and \(t_1=\exp(-2\pi i/\sqrt{e^*}d/\lambda)\). Here \(r\) is the reflection amplitude of a thick sample, \(t_1\) is the “pure” transmission amplitude, \(e^*\) is the complex dielectric permittivity of the sample, \(d\) is the sample thickness, and \(\lambda\) is the radiation wavelength. It has been assumed that the magnetic permeability \(\mu=1\) in the frequency range of our experiments. The transmittance for a two-layer system can be obtained in a similar way.\(^34,35\) Using Eq. (1), the absolute values of the complex conductivity \(\sigma = \sigma_1+i\sigma_2\) and dielectric permittivity \(\varepsilon = \varepsilon_1+i\varepsilon_2 = \sigma/i\varepsilon_0\omega\) can be determined directly from the measured spectra. Here \(\varepsilon_0\) and \(\omega=2\pi\nu\) are the permittivity of vacuum and the angular frequency, respectively. In general, the phase shift \(\phi_T\) in the transmittance experiment contains a possible ambiguity of \(2\pi n\), where \(n\) is an integer. To exclude this source of errors, the reflectance of a thick sample has been measured in addition to the transmittance experiments. A similar Mach-Zehnder arrangement allowed us to obtain the amplitude and the phase shift of the reflectance at the Verwey transition. In total, a set of four measured quantities was available to calculate the real and imaginary part of the dynamic conductivity in the THz frequency range.

In the infrared and visible frequency ranges \(30\) cm\(^{-1}\leq\nu<21\ 000\) cm\(^{-1}\), the complex conductivity has been obtained via the Kramers-Kronig analysis of the reflectivity of the thick sample. The reflectivity experiment were performed using Bruker IFS-113v and IFS 66v/S Fourier-transform spectrometers. Different sources, beam splitters, and optical windows allowed us to cover the complete frequency range.

In addition, the reflectance for the frequency range \(4\) cm\(^{-1}\leq\nu<40\) cm\(^{-1}\) has been calculated using the complex conductivity data, obtained by the transmittance technique described above. The combination of the results from two experimental techniques substantially expands the low-frequency limit of the available spectrum and the quality of the subsequent Kramers-Kronig transformation.

III. RESULTS AND DISCUSSION

Figure 1 shows the complex reflectance and transmittance of magnetite in the vicinity of the Verwey transition. All four quantities reveal strong changes at the transition temperature. The most dramatic changes are observed in the transmittance, which decreases by three orders of magnitude. Both transmittance and reflectance show a hysteretic behavior, characteristic for a first-order phase transition. However, the experimentally observed hysteresis extends over different temperature regimes: the hysteresis spans \(\Delta T\approx2\) K in the reflectance experiments, and is significantly less than 0.5 K in transmittance experiments. We suggest that this distinct difference probably results from the sample geometries, namely utilizing a thick sample (\(\approx5\) mm) in the reflectance and thin sample (\(\approx0.05\) mm) in the transmittance experiments. Except for the hysteresis region, all measurements revealed results coinciding within the experimental accuracy (Fig. 2). The effective overdetermination of the physical...
in Pr(Ca: Sr)MnO₃. We believe that the first-order character of the phase transition in magnetite interrupts the increase of the dielectric constant, observed at T < Tᵥ (dε/dT ≈ 1 K⁻¹ at low frequencies). Instead, the dielectric constant of magnetite abruptly jumps to negative values at the “metallic” side of the transition.

Figure 3 shows the frequency dependence of the conductivity and dielectric constant in magnetite. Symbols at low frequencies represent the results of the transmittance experiments. Solid lines above 30 cm⁻¹ have been obtained via the Kramers-Kronig analysis of the reflectance. The lower frame of Fig. 3 shows the frequency dependence of σₑ above and below the Verwey transition. The low-frequency conductivity in the insulating state is dominated by a power law in frequency (σₑ ≈ νˢ with s ~ 1.3) with a temperature-dependent amplitude and a weak temperature-dependent frequency exponent. As mentioned in Sec. I, a power-law behavior of the conductivity is a characteristic feature of hopping conduction between localized states.²³⁻²⁵ Previously, the frequency exponent s ~ 0.7 has been observed in magnetite at low temperatures and at kHz-frequencies,²¹ a value typical for the ac conductivity in the audio-frequency range. Substantially higher values of the exponent are expected at higher frequencies. Approaching the phonon-assisted-hopping regime, σₑ ≈ ν¹ has been predicted for low temperatures.³⁰ The transition to higher frequency exponents in the microwave frequency range has been discussed recently for doped semiconductors.⁴¹ The power-law term is generally referred to as the universal dielectric response²² and has been a recent matter of discussion concerning a possible universal super-linear power law with s ≥ 1 at high frequencies.⁴²

The frequency dependence of the dielectric constant in magnetite is shown in the upper panel of Fig. 3. The behavior of the dielectric constant in the insulating state agrees qualitatively with the power law of σₑ, i.e., σₑ increases with decreasing frequency. For s > 1 the low-frequency behavior can be approximated by εₑ(0) = εₑ(ν) ≈ ν⁻¹.

The dashed lines in Fig. 3 for the insulating state (T < Tᵥ) were obtained by fitting the experimental data to the expression

$$\sigma_e(\nu) = \sigma_{dc} + A \frac{(\nu/\nu_1)^s}{1 + (\nu/\nu_1)^4}.$$  

Here σₑ is the dc conductivity and A is the amplitude of the hopping process. Compared to the conventional form of this response²²⁻²⁵ we introduced an additional high-frequency cutoff [1 + (ν/ν₁)⁴]. The cutoff frequency ν₁ only weakly influences the conductivity in the low-frequency range but prevents the divergence of the spectral weight of σₑ ≈ ν¹ at high frequencies.

The analytical expression for the dielectric constant has been obtained by applying the Kramers-Kronig transformation⁴³ to Eq. (2), which gives the following analytical expression for the dielectric constant:

$$\epsilon_e(\nu) = \frac{\sigma_e(\nu) - \sigma_{dc}}{2\pi\nu\epsilon_0} \left[ \tan(\pi s/2) - \frac{(\nu/\nu_1)^s}{\cos(\pi s/2)} \sin \left( (s + 1) \frac{\pi}{4} \right) ight] + \left( \frac{\nu}{\nu_1} \right)^2 \cos \left( (s + 1) \frac{\pi}{4} \right).$$  

This expression is valid both for s < 1 and for s > 1.
To take into account the phononic and high-frequency electronic contributions, the sum of six Lorentzians has been added to expressions Eq. 1 and Eq. 2. Five Lorentzians at frequencies of 106, 152, 247, 361, and 574 cm\(^{-1}\) represent phonon contributions. An additional overdamped oscillator at 4270 cm\(^{-1}\) approximates the first excitation of the electronic origin. The given characteristic frequencies agree well with the published data.\(^{13-15}\) In order to simplify the analysis of the THz conductivity and obtain the essential physics of the problem, the contributions from the Lorentzians were fixed for all three temperatures in Fig. 3. The parameters obtained from simultaneous fitting of the conductivity and dielectric constant are given in Table I.

A remarkable feature of the parameters for the low-temperature insulating phase in Table I is the low value of the cutoff frequency \(\nu_1\). Indeed, a qualitative examination of the conductivity frame of Fig. 3 reveals that the hopping process \(\sigma_1 \sim \nu^s\) and the low-frequency edge of the electronic excitation \((\nu > 1000 \text{ cm}^{-1})\) are well separated in energy. This indicates that the cutoff frequency of the hopping must be in the range \(\sim 100 \text{ cm}^{-1}\), in agreement with the fits. This result is rather surprising and suggests that the low-temperature conductivity is governed by phonons.

In contrast to the gradual variation of \(\varepsilon_1(\omega, T)\) and \(\sigma_1(\omega, T)\) at low temperatures, dramatic changes are observed at the Verwey transition. At low frequencies the conductivity increases by more than one order of magnitude. This increase is substantially smaller than a change of two orders of magnitude in the dc conductivity (see insets of Fig. 2). This difference is the result of a power-law frequency dependence of the conductivity in the insulating state. As a consequence, for frequencies close to 35 cm\(^{-1}\) we observe almost no conductivity change at \(T_V\). This observation has been verified in a separate temperature-dependent experiment at \(\nu = 35 \text{ cm}^{-1}\) (not shown).

As discussed above, the decrease of the dielectric constant at the transition to the conducting state is not expected within the scope of a conventional metal-insulator transition scenario. In order to find an explanation to this experimentally observed behavior, we mention that a negative dielectric constant was observed in some experiments.

![Graph](https://i.imgur.com/5.png)

**FIG. 3.** (Color) Frequency dependence of the conductivity \((\sigma_1, \text{lower panel})\) and the dielectric constant \((\varepsilon_1, \text{upper panel})\) of magnetite above and below \(T_V = 123 \text{ K}\). Symbols below 40 cm\(^{-1}\) and solid lines represent the experimental data; dashed lines are the model calculations. Experimental data have been obtained from transmittance and phase shift of a thin Fe\(_3\)O\(_4\) plate below 40 cm\(^{-1}\), and via the Kramers-Kronig analysis of the reflectance of the thick sample above 30 cm\(^{-1}\). The main contribution to the model calculations is given by the hopping term \([\sigma^* \sim \nu^s, \text{Eq. (2)}]\) below \(T_V\) and a Drude term [Eq. (4)] above \(T_V\).

**TABLE I.** Parameters of the THz-frequency excitations in magnetite close to the Verwey transition that correspond to Eqs. (2)–(4).

<table>
<thead>
<tr>
<th>(T (K))</th>
<th>(\nu_1 (\text{cm}^{-1}))</th>
<th>(s)</th>
<th>(A (\Omega^{-1} \text{ cm}^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>100</td>
<td>70</td>
<td>1.5</td>
<td>39</td>
</tr>
<tr>
<td>120</td>
<td>60</td>
<td>1.2</td>
<td>44</td>
</tr>
<tr>
<td>(T_{1/2 \pi \tau} (K))</td>
<td>(\sigma_{dc} (\Omega^{-1} \text{ cm}^{-1}))</td>
<td></td>
<td></td>
</tr>
<tr>
<td>130</td>
<td>40</td>
<td>37</td>
<td></td>
</tr>
</tbody>
</table>
stant naturally follows by assuming a Drude response of quasi-free carriers,
\[ \sigma_{\text{DR}}^* = \sigma_{\text{dc}}/(1 - i2\pi\nu\tau), \]
where \( \sigma_{\text{dc}} \) and \( 1/2\pi\tau \) are dc conductivity and the scattering rate, respectively. In addition to the negative changes in \( \varepsilon_1 \), the conductivity in the metallic state shows a slight downward curvature in the frequency dependence (lower panel of Fig. 3). This provides a further argument to include a Drude term to the model. Indeed, the simplest way to reproduce the frequency of the coherent process, its spectral weight, \( S = \frac{ne^2}{m_{\text{eff}}} \exp\left(-\frac{\Delta_m}{k_B T}\right) \),
\[ \text{(6)} \]
where \( n, e, \) and \( m_{\text{eff}} \) are concentration, charge, and effective mass of the charge carriers, respectively. In this expression the effective concentration of charge carriers is temperature activated and is governed by an energy gap \( \Delta_m \). The estimate of the effective mass \( m_{\text{eff}} \approx 10^5 m \) has been obtained from the infrared conductivity\(^{14} \) and from the small-polaron analysis of photoemission of the same sample.\(^{11} \) With this estimate for \( m_{\text{eff}} \) the observed small spectral weight of the Drude process corresponds to \( \Delta_m \approx 40 \text{ meV} \), which correlates with a weakly activated dc conductivity just above \( T_V \) (right inset in Fig. 2). Within the same picture, the Verwey transition in magnetite corresponds just to an increase of the characteristic activation energy at the insulating side of the transition. The drop of dc conductivity at the transition can be ascribed to a gap change from \( \Delta_m \approx 40 \text{ meV} \) (conducting state) to \( \Delta_i \approx 90 \text{ meV} \) (insulating state). This new value of the energy gap agrees with the activation energy of dc conductivity\(^{16-20} \), thermolectric power,\(^{12} \) and with the gap values obtained in photoemission experiments.\(^{9-11} \) The presented description implies the interpretation of the Verwey transition as an insulator-semiconductor transition with a change in activation energy induced by a crystal structure or by electronic configuration.

IV. CONCLUSIONS

Terahertz and infrared conductivity in magnetite has been obtained on both sides of the Verwey metal-to-insulator transition. Two orders of magnitude change in dc conductivity halves at millimeter frequencies and even disappear for \( \nu \approx 30 \text{ cm}^{-1} \). In the far-infrared range, the effect of the transition remains solely in changing the dielectric constant \( (\varepsilon_1) \) of magnetite. The change of \( \varepsilon_1 \) during the transition into the conducting state is negative, which is in surprising contrast to an expected (positive) divergence of the dielectric constant at a metal-insulator transition. Within the simple model analysis the conductivity mechanism switches between the hopping of localized carriers below \( T_V \) and itinerant motion above \( T_V \). These results evidence the formation of the coherent state on the metallic side of the Verwey transition with high mobility of the charge carriers. Together with the recent photoemission data on the same samples, the interpretation of the Verwey transition as an insulator-semiconductor transition is suggested.

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