

Electrically induced phase transition in α -(BEDT-TTF)₂I₃: Indications for Dirac-like hot charge carriers

T. Peterseim, T. Ivek, D. Schweitzer, and M. Dressel

I. Physikalisches Institut, Universität Stuttgart, Pfaffenwaldring 57, 70550 Stuttgart, Germany

(Received 11 April 2016; revised manuscript received 26 May 2016; published 14 June 2016)

The two-dimensional organic conductor α -(BEDT-TTF)₂I₃ undergoes a metal-insulator transition at $T_{CO} = 135$ K due to electronic charge ordering. We have conducted time-resolved investigations of its electronic properties in order to explore the field- and temperature-dependent dynamics. At a certain threshold field, the system switches from a low-conducting to a high-conducting state, accompanied by a negative differential resistance. Our time-dependent infrared investigations indicate that close to T_{CO} , the strong electric field pushes the crystal into a metallic state with optical properties similar to the one for $T > T_{CO}$. Well into the insulating state, however, at $T = 80$ K, the spectral response evidences a completely different electronically induced high-conducting state. Applying a two-state model of hot electrons explains the observations by excitation of charge carriers with a high mobility. They resemble the Dirac-like charge carriers with a linear dispersion of the electronic bands found in α -(BEDT-TTF)₂I₃ at high pressure. Extensive numerical simulations quantitatively reproduce our experimental findings in all details.

DOI: [10.1103/PhysRevB.93.245133](https://doi.org/10.1103/PhysRevB.93.245133)

I. INTRODUCTION

The charge-transfer salt α -(BEDT-TTF)₂I₃ is probably the best studied model compound of a metal-insulator transition in two-dimensional electron systems [1,2] that is supposed to be driven by electronic charge order [3–5]. From NMR [6,7] and optical spectroscopy [8–13] as well as x-ray-diffraction studies [14], it is well known that below $T_{CO} = 135$ K charge disproportionation develops on the BEDT-TTF molecules. Since its discovery three decades ago, interest in the title compound has never faded because it exhibits a rich temperature-pressure phase diagram, with a number of intriguing quantum phenomena ranging from electronic ferroelectricity [15–17] to superconductivity [18,19], from nonlinear transport [20,21] to zero-gap semiconductivity [22,23] characterized by Dirac cones and massless Dirac fermions [24–27], but also the appearance of persistent photoconduction [28], photoinduced phase transition [29–31], and nonlinear ultrafast optical response [32].

At room temperature, α -(BEDT-TTF)₂I₃ consists of layers formed by the BEDT-TTF molecules that are separated along the c' direction by I₃[−] anions creating an electronically two-dimensional system, as displayed in Fig. 1. At $T_{CO} = 135$ K, the symmetry class of the crystal changes from $P\bar{1}$ to $P1$ at low temperatures, thus breaking the inversion symmetry between the molecules A and A'. Now, all four molecules are crystallographically independent, and each molecule in the unit cell carries a distinct charge different from the others. As shown in Fig. 1(c), the charge order forms horizontal stripes with molecule B still being charge-rich and molecule C charge-poor, but either molecule A or A' being charge-poor/rich. The charge ordering leads to a ferroelectric state that was probed by second-harmonic generation of light [32–34] as a direct proof of the loss of the inversion symmetry. These studies have also shown that different ferroelectric domains exist in a single crystal separated from each other by domain walls. Indications of an influence of cooling rate and pressure on the amount of domains and their size were found, although a systematic study is still missing and necessary.

Recent dielectric measurements [12,15] detected two modes with a dielectric strength $\Delta\epsilon$ between 400 and 5000 for the a and b direction, indicating a strong ferroelectric response with no temperature dependence below T_{CO} . The first mode is present at all temperatures and is ascribed to a charge-density wave, which is screened by activated charge carriers since the relaxation time follows the temperature dependence of the dc conductivity. The second mode appears upon further cooling below $T = 75$ K. It is suggested that this response is caused by solitons and domain walls that are fixed to pinning centers. Alternatively, the existence of polar and nonpolar stacks of the BEDT-TTF molecules was suggested [17], preventing long-range ferroelectricity.

In a complementary approach, the electrodynamic properties of α -(BEDT-TTF)₂I₃ at the metal-insulator phase transition have been explored by nonlinear and time-dependent studies [35,36]. Particular attention was drawn by the observation of photoinduced phase-transition phenomena [29,30,37], photoconductivity [28,38] connected with a memory effect [39,40], nonlinear conductivity [2,21,41], and zero-gap states under pressure [23,27,42,43]. Here we focus on the electrically induced phase transition in α -(BEDT-TTF)₂I₃ by investigating the time-resolved transport and optical properties. We then apply a model of nonequilibrium charge carriers to our observations, explaining our findings in all details.

II. EXPERIMENTAL RESULTS

The time-, temperature-, and field-dependent transport and optical studies have been conducted on α -(BEDT-TTF)₂I₃ single crystals, as grown by electrochemical methods [1,44]. The experimental details are given in the Supplemental Material [45].

In addition to conventional dc transport measurements in four-point geometry (cf. Figs. S1 and S2), dynamical resistivity studies have been performed by applying a voltage of U_S across the single crystal of α -(BEDT-TTF)₂I₃ for a few milliseconds and measuring the total current flow I_{tot} as a function of

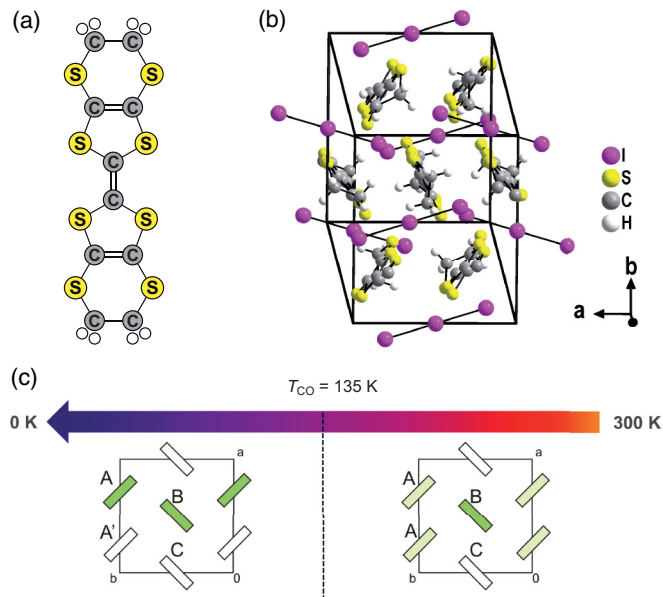


FIG. 1. (a) Sketch of the bis-(ethylenedithio)tetrathiafulvalene molecule, called BEDT-TTF. (b) Drawing of the α -(BEDT-TTF)₂I₃ unit cell with orientation along the c' direction. The unit cell contains four BEDT-TTF and two I₃ molecules. The cations are arranged in a herringbone-like structure within the ab plane separated from each other along the c' direction by the anions. (c) Illustration of the development of the charge disproportionation and crystal symmetry in α -(BEDT-TTF)₂I₃ when cooling from room temperature to 0 K. The ab plane is displayed with the four BEDT-TTF molecules labeled by A, B, and C; the anions are not included for clarity reasons. The A molecules are related to each other by a point inversion on their connecting line. The opacity of the green color represents the charge imbalance between the molecules. At $T = 300$ K, there is an imbalance between molecules B and C. Below T_{CO} , the point inversion is lost and the A molecules become inequivalent. A stripelike charge-order pattern develops along the b direction.

time. In the metallic but also well into the insulating state, the response is basically constant at low voltage, indicating a steady state. In Fig. 2(b) it can be seen, however, that for $U_S = 80$ V applied at low temperatures ($T = 80$ K, for instance), heating continuously increases the current until the sample suddenly turns metallic at a certain delay time τ_d . In a first step (Sec. II A), we want to characterize how this switching evolves with time and how the behavior depends on temperature and electric field applied. In Sec. II B we will then present optical investigations that will elucidate the nature of the electrically induced metallic state.

A. Transport studies

In Fig. 3, the time dependence of the sample resistance is plotted as a function of the applied electric field for two selected temperatures. At $T = 125$ K, the sample resistance R_{sample} is approximately 19 k Ω . In the time frame under inspection, it jumps abruptly from the insulating state into a high-conducting state above a certain threshold field that is marked by the color change from green to blue in Fig. 3. The resistance is reduced by two orders of magnitude to

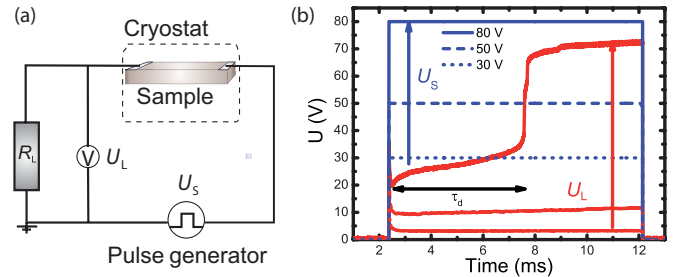


FIG. 2. (a) For time-dependent transport studies, a voltage pulse U_S of some milliseconds duration is applied; the voltage drop U_L across a load resistor measures the total current $I_{\text{tot}} = U_L/R_L$. (b) Time dependence of the source voltage U_S (blue lines) and drop U_L across the load resistor (red curves) measured at $T = 80$ K along the a direction of α -(BEDT-TTF)₂I₃. After a short switching pulse, a steady-state regime is reached for low voltages (dashed and dotted lines). Above a certain threshold field, however, the sample switches from a low-conducting, insulating state into a high-conducting state, which is marked by a steep increase of the current.

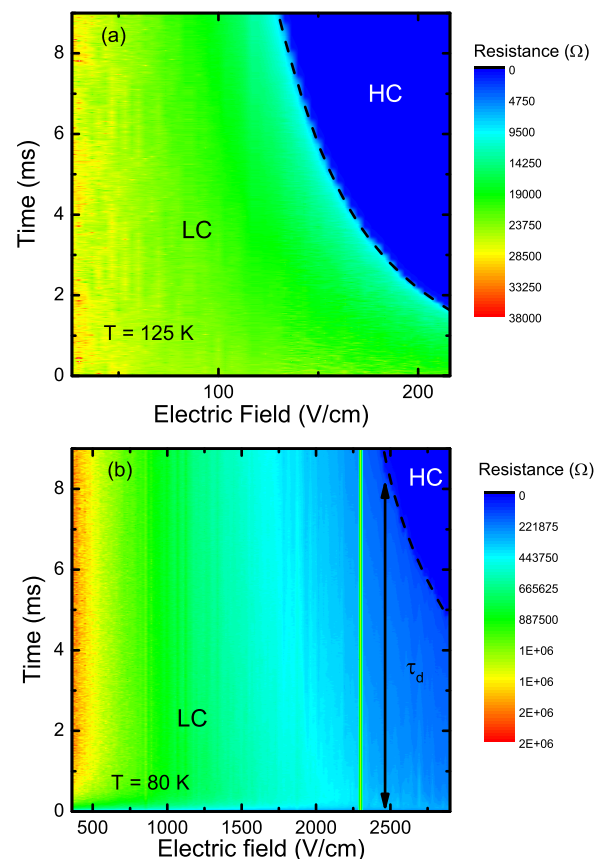


FIG. 3. Contour plot of the time-dependent sample resistance of α -(BEDT-TTF)₂I₃ in a time frame of 10 ms for different electric fields E_{sample} recorded at (a) a temperature close to the charge-order transition, $T = 125$ K, and (b) well below, $T = 80$ K. The blue area corresponds to a metallic behavior (high conducting, HC), while the green to red areas indicate the insulating state (low conducting, LC). With increasing field, the delay time τ_d to enter the metallic state decreases. Note that in both frames different scales have been used for the applied field and for the observed resistance, respectively.

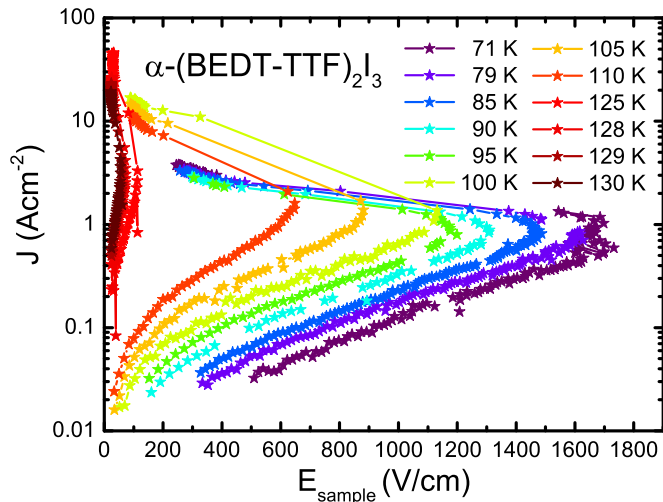


FIG. 4. Current density J as a function of the electric field E_{sample} recorded at a α -(BEDT-TTF) $_2$ I $_3$ single crystal at different temperatures between $T = 130$ and 80 K as indicated. At all temperatures, a negative differential regime is observed, where the threshold electric field increases with decreasing temperature.

$R_{\text{sample}} \approx 200 \Omega$. For the contour plot, it becomes clear that the switching takes place with a certain delay time τ_d , which decreases with increasing electric field E_{sample} in a nonlinear manner.

For lower temperatures, $T = 80$ K, the behavior is rather similar; however the total resistance $R_{\text{sample}} \approx 1 \text{ M}\Omega$ is much higher and decreases slightly with increasing voltage due to the nonlinear behavior of the contact resistance [21]. Note that at reduced temperatures the electric field must be significantly increased to induce the high conducting state. It should also be pointed out that despite the different absolute values of resistivity recorded, the step height between the low- and high-conducting range is again about two orders of magnitude, as in the case of $T = 125$ K.

The calculated current-voltage characteristic, J versus E_{sample} , is displayed in Fig. 4 for the temperature range from $T = 70$ to 130 K. The J curves exhibit a regime of a negative differential resistance (NDR), where the current jumps to a much larger value, while the electric field at the sample E_{sample} drops significantly. This curve reveals a so-called S-shape (current-controlled), in contrast to an N-curve (voltage-controlled), where the current drops to smaller values with increasing electric field due to a reduction of the carrier mobility. The NDR appears above a certain threshold field E_{thresh} , which increases with lowering T . Before the NDR regime, the current density increases linearly reflecting Ohmic behavior. In our context, this means that the conductivity must be significantly enhanced by generating high-mobility carriers.

From Fig. 4, we extract the threshold electric field E_{thresh} and the threshold current density J_{thresh} , and we plot the parameter as a function of temperature in Fig. 5(a). E_{thresh} decreases linearly with T up to T_{CO} where it vanishes, as expected; there is no NDR observed in the metallic state above T_{CO} . This agrees with measurements in the current mode performed in α -(BEDT-TTF) $_2$ I $_3$ below 100 K [41,46]; in other organic materials, similar observations have been reported

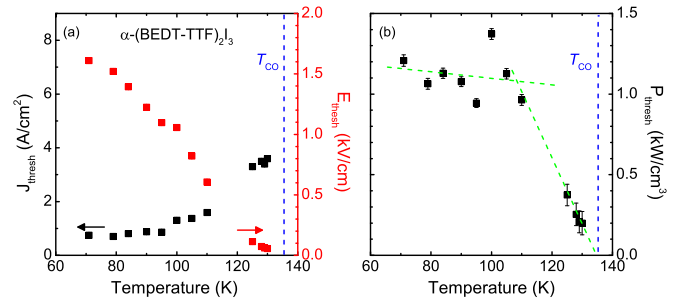


FIG. 5. (a) Temperature dependence of the threshold current density J_{thresh} (black, left axis) and threshold electric field E_{thresh} (red, right axis). (b) Power density P_{thresh} below the transition temperature T_{CO} of α -(BEDT-TTF) $_2$ I $_3$. The green dotted lines are guides to the eye. The blue dashed vertical lines mark the metal-insulator transition temperature T_{CO} .

[47]. There, the threshold field E_{thresh} also rises continuously as T is reduced.

Let us compare the absolute value of the NDR threshold fields found in our experiments to other systems. In the case of typical charge-density-wave materials [48], threshold fields below 1 V/cm are typically observed, while the electric field values we extract from α -(BEDT-TTF) $_2$ I $_3$ are several orders of magnitude larger. On the other hand, avalanche effects or dielectric breakdown are found in semiconductors or insulators only for fields exceeding 1 MV/cm . However, the observed temperature dependence is also distinct from those phenomena. For a charge-density wave, the threshold field follows a $E_{\text{thresh}} \propto \exp\{-T/T_0\}$ behavior [49], in contrast to dielectric breakdown, where it behaves like $E_{\text{thresh}} \propto \exp\{T_0/T\}$. The linear T dependence observed here in α -(BEDT-TTF) $_2$ I $_3$ does not follow either of them. Hence we conclude that the nonlinear transport behavior in charge-ordered materials plays a special role and cannot be simply classified within the previous mentioned phenomena.

In contrast to E_{thresh} , the current density $J_{\text{thresh}}(T)$ does not follow a linear temperature behavior; it seems to diverge toward T_{CO} and approaches a constant value at low T . Similar results have been reported previously [46]. From both quantities, the power density at the threshold, $P_{\text{thresh}} = E_{\text{thresh}} J_{\text{thresh}}$, can be readily calculated [Fig. 5(b)]. Interestingly, the power necessary to switch the conduction state rises steeply as the temperature drops below T_{CO} ; only for $T < 120$ K does P_{thresh} remain basically constant. This temperature dependence resembles the behavior of the resistivity plotted in Fig. S2, where the largest modification occurs in a narrow temperature range $120 < T < 135$ K where the energy gap opens.

To ensure that this effect is not caused by Joule heating, we have conducted a thorough calculation of the stored energy and the possible temperature rise; details are given in the Supplemental Material [45]. Even under the assumption that 100% of the electrical power is used for heating the crystal, the experiments performed at $T = 80$ K will never lead to sample temperatures above 96 K. At least far below $T_{\text{CO}} = 135$ K, we can therefore exclude that the switching from the low-conducting to the high-conducting state is caused by Joule heating.

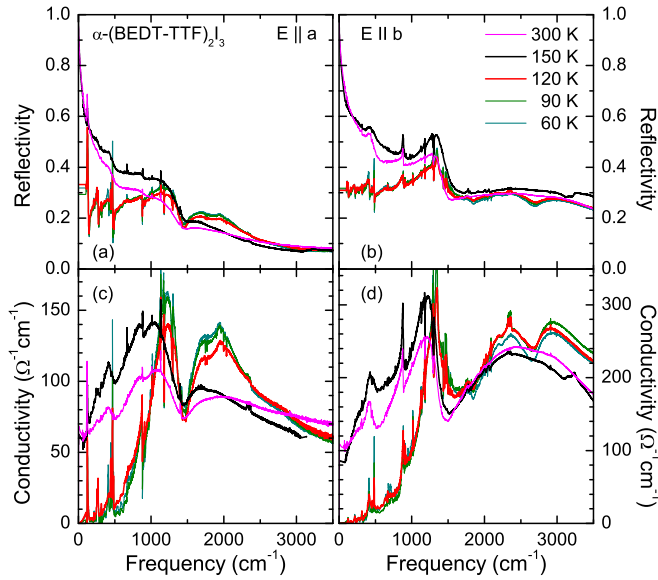


FIG. 6. Optical properties of α -(BEDT-TTF) $_2$ I $_3$ for different temperatures as indicated. The upper panels (a) and (b) show the reflectivity for the polarization of the electric field $E \parallel a$ and $E \parallel b$. The corresponding optical conductivity is plotted in the lower panels (c) and (d); note the different vertical scales.

B. Optical studies

Beyond these time-dependent transport studies, more insight into the nature of the electronic state can be obtained from the frequency-dependent conductivity [50]. It is well known that the metal-insulator transition of α -(BEDT-TTF) $_2$ I $_3$ at $T_{CO} = 135$ K has a strong influence on the optical response probed by polarized infrared reflection measurements. As illustrated in Fig. 6, the reflectivity decreases below approximately 1500 cm^{-1} when the temperature drops below T_{CO} , while $R(\omega)$ increases in the midinfrared region. The corresponding optical conductivity is plotted in Figs. 6(c) and 6(d), demonstrating the shift in spectral weight from the frequency range below the gap at approximately 600 cm^{-1} to the midinfrared peak around 200 cm^{-1} [51]. The change takes place within a rather narrow temperature interval between $T = 140$ and 120 K with basically no variation below.

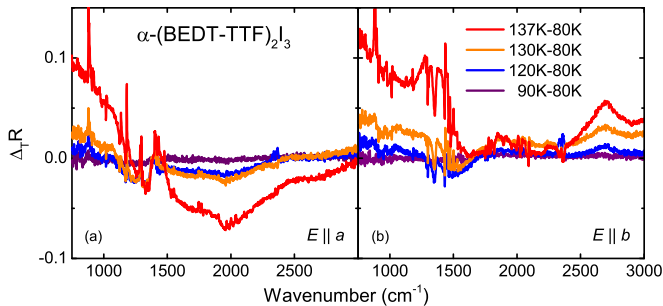


FIG. 7. Difference of the optical reflectivity at $T = 137$ K (red), 130 K (orange), 120 K (blue), and 90 K (purple) with respect to the reflectivity at $T = 80$ K. Panels (a) and (b) correspond to the polarizations $E \parallel a$ and $E \parallel b$, respectively.

Since we are interested in how the reflectivity changes in certain spectral regions, in Fig. 7 we plot the difference in reflectivity $\Delta_T R(\nu)$ between various temperatures above and below T_{CO} . When comparing $T = 120$ and 80 K, for instance, $\Delta_T R$ exhibits almost no modification besides a slight shift of the energy gap in the charge-ordered phase. Pronounced spectral changes, however, are found between the two phases, i.e., looking at $T = 137$ and 80 K. When entering the charge-ordered state below T_{CO} , the low-frequency reflectivity rises on the expense of the midinfrared region. No significant differences are observed between the two polarizations $E \parallel a$ and $E \parallel b$.

To characterize the electric-field-induced metallic state spectroscopically and to draw a more complete picture of the dynamics at the metal-insulator transition of α -(BEDT-TTF) $_2$ I $_3$, we have performed time-resolved infrared measurements by employing step-scan Fourier-transform infrared spectrometer Bruker Vertex 80v; for technical details, see Ref. [52]. The time resolution was set to $20 \mu\text{s}$, and 1000 time steps were recorded for 20 ms. The data were acquired 1 ms before the voltage pulse was initialized. To improve the signal-to-noise ratio, we averaged over 20 scans. The spectral resolution was 4 cm^{-1} and the recorded frequency ranged from $\nu = 800$ to 4000 cm^{-1} . The time-resolved measurements were performed in the ac mode, highlighting the changes in reflectivity $\Delta_T R(\nu, t)$ as time t elapses, where $\Delta_T R(\nu, t) = R(\nu, t) - R(\nu, 0)$. It is of interest to compare this with the change of reflectivity when the temperature T is varied with respect to a reference temperature T_0 : $\Delta_T R(\nu, T) = R(\nu, T) - R(\nu, T_0)$, as plotted in Fig. 7.

1. Close to metal-insulator transition: $T = 125$ K

In Fig. 8, the time- and frequency-dependent change of the reflectivity $\Delta_T R(\nu, t)$ at $T = 125$ K is plotted as recorded for the a and b direction when an electric field of $E_{\text{sample}} = 216 \text{ V/cm}$ is applied parallel to the crystallographic a axis. Reflectivity data $R(\nu, t)$ in the midinfrared range are acquired during a period of 20 ms for both directions. After 1 ms, the voltage pulse is applied to the sample, but it takes another few milliseconds before the reflectivity changes appreciably. Only 4 ms after the voltage pulse has started does $R(\nu, t)$ increase for $\nu < 1500 \text{ cm}^{-1}$, which is accompanied by a drop of the signal between 1500 and 2500 cm^{-1} . Qualitatively the behavior is similar for both directions. After the voltage pulse has stopped at $t = 11$ ms, the signal keeps increasing almost to the end of the measurement window. The influence of the electric field on the material exceeds the pulse duration; any modification remains longer than the recorded time frame.

To analyze the time dependence of the reflectivity $R(\nu, t)$ in more detail, we take spectral and time slices at the position marked in Fig. 8 by black arrows. In panels (a) and (b) of Fig. 9, the evolution of the $\Delta_T R(\nu, t)$ spectra is shown for six different points in time, ranging from $t = 0.5$ to 18 ms. Before the pulse is applied (at $t = 0.5$ ms), the reflectivity is steady. Immediately following the applied field, the reflectivity rises, but at 2 ms $\Delta_T R(\nu, t)$ comes to a halt at a minuscule level, where it stays until $t = 4$ ms. From our time-resolved transport measurements presented in Fig. 3, we can extract that at $T = 125$ K the electrically induced switching occurs

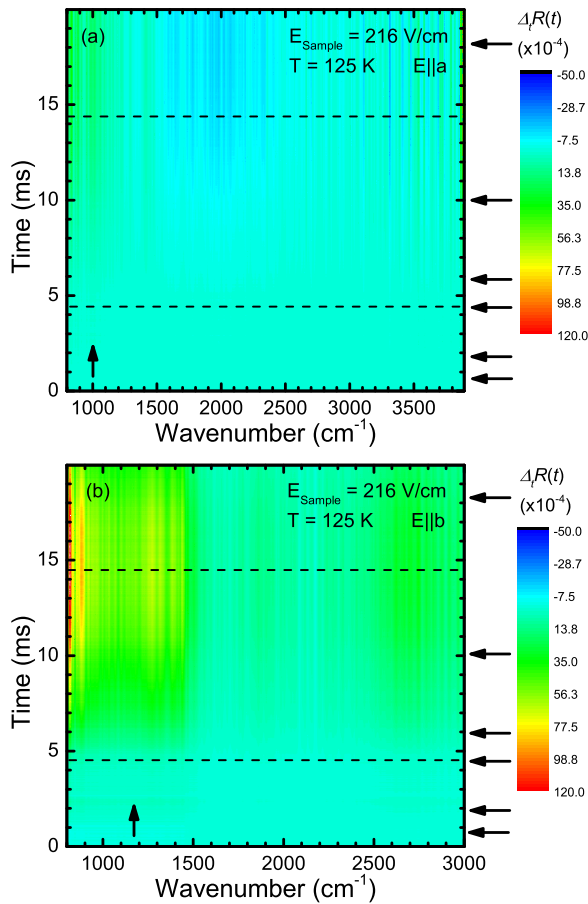


FIG. 8. Contour plot of the reflectivity change $\Delta_T R(\nu, t)$ along the (a) a and (b) b directions of α -(BEDT-TTF) $_2$ I $_3$ at $T = 125$ K after applying an electric field of $E_{\text{sample}} = 216$ V/cm along the a axis lasting for 10 ms. The dashed horizontal lines mark the onset and end of the voltage pulse, whereas the arrows mark the position where the spectra and the time slices were extracted from the frequency- and time-dependent spectrum, and they are displayed in Fig. 9. The vertical stripes in the spectrum are due to instabilities of the interferometer mirror during the step-scan run.

1.7 ms after the field is applied. Thus we cannot relate the phase transition detected in the resistivity drop with the change of the reflectivity at 5 ms. We conclude that two different effects are responsible for these observations. By taking a closer look at the spectral shape of $\Delta R(\nu, t)$ after 5 ms and comparing them with the reflectivity difference $\Delta_T R$ presented in Fig. 7, we immediately recognize the similarity in the difference spectra $\Delta_T R(\nu, T)$ between $T = 137$ and 80 K. In other words, the electric pulse transforms the crystal into the metallic phase after 4 ms.

These observations and conclusions are confirmed when we analyze the time-dependent behavior for two selected frequencies, $\nu = 1000$ and 1354 cm^{-1} , plotted in Figs. 9(c) and 9(d). The signal starts to rise slightly with the onset of the voltage pulse, but it soon levels off. The electrically induced switching point at $t = 2.7$ ms is not reflected in the $\Delta_T R(t)$ spectra. Only 4 ms after the voltage is applied does the reflectivity start to rise linearly. This continues well beyond the end point of the pulse; $\Delta_T R(t)$ saturates at about 15 ms

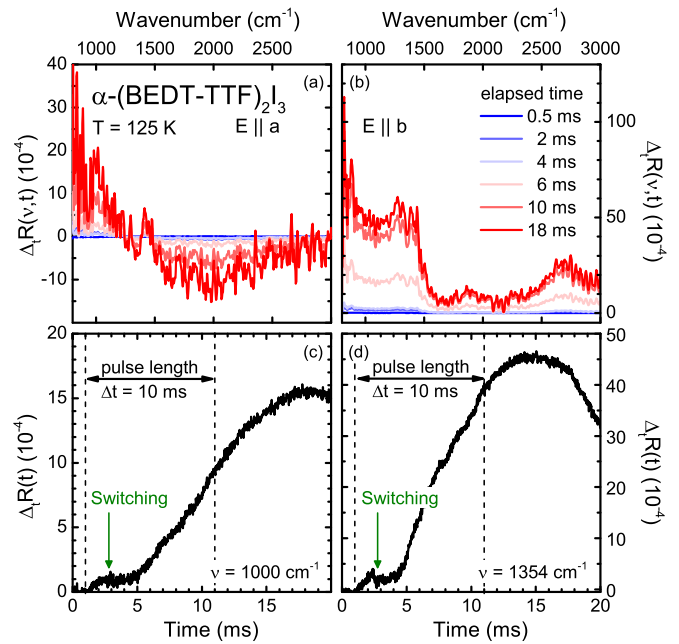


FIG. 9. (a,b) Variation of the reflectivity $\Delta_T R(\nu, t)$ spectra of α -(BEDT-TTF) $_2$ I $_3$ for different times at $T = 125$ K. The data in the frequency range from $\nu = 800$ to 3000 cm^{-1} are extracted from Fig. 8 for both polarizations (a) $E \parallel a$ and (b) $E \parallel b$. The lower panels display the time profile of the reflectivity $R(\nu, t)$ taken at selected frequencies (c) $\nu = 1000$ cm^{-1} and (d) 1354 cm^{-1} between 0 and 20 ms. When the electric field is applied, $R(\nu, t)$ increases only slightly and then remains constant in the high-conducting state. With a delay of a few milliseconds, the signal rises linearly until it saturates after the voltage pulse is switched off; the reflectivity then decays linearly. The vertical dotted lines mark the start and end points of the voltage pulse.

and then decreases linearly. With slight variations, the overall behavior is similar for both polarization directions.

Several peculiarities call for further discussion:

(i) We suggest that in the first 4 ms of the voltage pulse, a fraction of the sample is transferred into a metallic phase. This picture is supported by spatially resolved Raman measurements of a current-induced phase transition under steady-state conditions [53]. Mori *et al.* observed a modification of the spectral intensity of an electron-molecular vibrational (emv)-coupled mode of the BEDT-TTF molecules, which they ascribed to the variation of the electronic background and therefore to an electrically induced metallic state. In contrast, a temperature-sensitive vibrational mode of the I $_3^-$ anions does not reveal any modification. Note, however, that measurements under steady current flow are likely to be subject to heating [54]. Since we cannot correlate the observed switching from a low- to high-conducting state to the temporal dynamics of the sample resistivity, we interpret the change of reflectivity by a thermally induced phase transition of a few spatially limited areas. The electrical current is affected by some inhomogeneously distributed impurities and cracks within the sample, leading to local heating.

(ii) It is a remarkable fact that the sample is already in the highly conducting (metallic-like) state according to the transport measurements, but the reflectivity changes only after

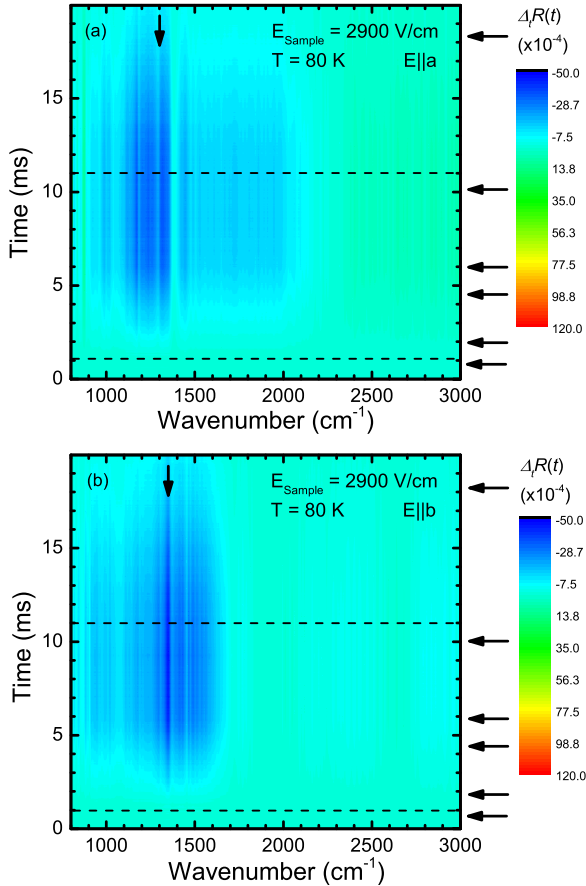


FIG. 10. Contour plot of the reflectivity change $\Delta_I R(\nu, t)$ of α -(BEDT-TTF) $_2$ I $_3$ for both polarizations at $T = 80$ K after applying an electric field of $E_{\text{sample}} = 2900$ V/cm along the a axis lasting for 10 ms. The same color code is used as in Fig. 8. The dashed horizontal lines mark the onset and end of the voltage pulse, whereas the arrows mark the position where the spectra and the time slices were extracted from the frequency- and time-dependent spectrum, and they are displayed in Fig. 11.

$t = 5$ ms and grows further until 15 ms, although the pulse is already switched off at $t = 11$ ms. We suggest that the system is in a highly conducting state due to an excited electronic system. These “excited” or “hot” charge carriers couple to the lattice subsystem to which the energy is eventually transferred with a certain delay time. This leads to a heating-up of the sample across the phase transition causing the reflectivity change. Calculations of the heating effects confirm this picture [45]. Here we assumed that the entire electric energy provided by the voltage pulse is completely converted into heat, and the sample temperature exceeds T_{CO} .

(iii) The excited charge carriers relax back to their initial ground state by releasing the energy to the crystal lattice. $\Delta_I R(\nu, t)$ saturates when the energy transferred to the phonon bath is balanced by the outflow to the heat sink of the sample holder. Eventually, all energy is stored in the lattice subsystem; without any influx, the temperature rises leading to a linear drop of the reflected signal. This picture consistently explains also our electric pump-probe experiments [21] where the resistivity recovers in an exponential manner within a few

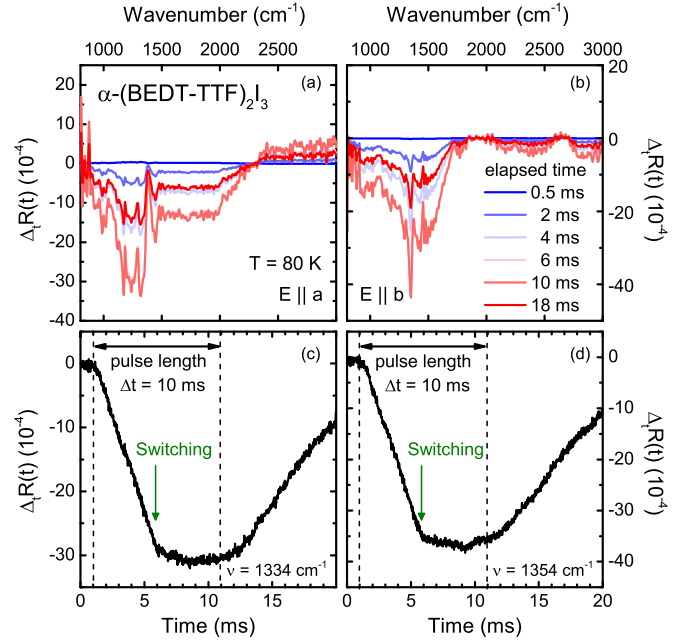


FIG. 11. (a,b) Variation of the reflectivity $\Delta_I R(\nu, t)$ spectra of α -(BEDT-TTF) $_2$ I $_3$ for different times at $T = 80$ K. The data in the frequency range from $\nu = 800$ to 3000 cm^{-1} are extracted from Fig. 10 for both polarizations (a) $E \parallel a$ and (b) $E \parallel b$. The lower panels display the time profile of the reflectivity $R(\nu, t)$ taken at selected frequencies (c) $\nu = 1000$ cm^{-1} and (d) $\nu = 1354$ cm^{-1} between 0 and 20 ms. When the electric field is applied, $R(\nu, t)$ increases only slightly and then remains steady in the high-conducting state. With a delay of a few milliseconds, the signal rises linearly until it saturates after the voltage pulse is switched off; the reflectivity then decays linearly. The vertical dotted lines mark the start and end points of the voltage pulse.

milliseconds after the highly conducting state was initialized by a 3-ms-long voltage pulse.

(iv) The linear decrease in $\Delta_I R(\nu, t)$ after 17 ms can be quantitatively described by Newton’s law of cooling, $Q_{\text{cool}} = -\lambda_{\text{therm}}(T_L - T_0)$, where λ_{therm} is the thermal conductivity, T_0 denotes the environment temperature, and T_L is the temperature of the heated lattice. Due to the elevated temperature, thermal radiation was neglected as a relevant factor.

2. Well below the metal-insulator transition: $T = 80$ K

Now, we consider the switching behavior of α -(BEDT-TTF) $_2$ I $_3$ at $T = 80$ K, i.e., at temperatures far below T_{CO} . The frequency- and time-dependent spectra $\Delta_I R(\nu, t)$ are displayed in Fig. 10 for the polarizations $E \parallel a$ and $E \parallel b$, under an electric field strength $E_{\text{sample}} = 2900$ V/cm. The voltage pulse arrives at the sample at time 1 ms and lasts for 10 ms. In both spectra, a change of $\Delta_I R(\nu, t)$ can be recognized right with the onset of the pulse. The important point is, however, that in contrast to the behavior at $T = 125$ K depicted in Fig. 8, the reflectivity gets smaller and thus $\Delta_I R(\nu, t)$ is mainly negative. It persists until the end of the record time, although it relaxes back about 4 ms after the voltage pulse has been switched off.

In Fig. 11 the frequency-dependent $\Delta_I R(\nu, t)$ is plotted for different points in times, extracted as slices from Fig. 10.

Immediately after the voltage pulse has arrived at 1 ms, the negative signal increases linearly, saturates at $t \approx 6$ ms, and eventually drops after 12.5 ms. For $E \parallel a$ there seems to be one isosbestic point around 2300 cm^{-1} ; above that frequency $\Delta_t R(\nu, t)$ develops similarly, but with opposite, i.e., positive sign. For $E \parallel b$ two frequencies can be identified, $\nu \approx 1900$ and 2650 cm^{-1} , with basically no temporal development, but the changes $\Delta_t R(\nu, t)$ are always negative. The spectra differ significantly from those recorded at $T = 125 \text{ K}$ (Fig. 9) and are interpreted as a reduction of the electronic band and the emv-coupled modes in the case of the a and b direction, respectively. The comparison with the temperature-modulated reflectivity $\Delta_T R(\nu, T)$ shown in Fig. 7 reveals a small increase of the sample temperature of only a few Kelvin, for instance from $T = 80$ to 90 K . Since we do not observe a positive signal below 1000 cm^{-1} , the energy gap does not shrink, in excellent agreement with our estimation of an upper limit of the sample temperature caused by Joule heating [45], which predicts a maximum temperature of 96.5 K in the high-conducting state, which is still far below $T_{\text{CO}} = 135 \text{ K}$. Therefore, we believe that for our 80-K experiments, no thermally induced phase transition is responsible for the switching from the low-conducting to the high-conducting state.

The time evolution of $\Delta_t R(\nu, t)$ at two frequencies is displayed Figs. 11(c) and 11(d). The signal decreases in a linear fashion right after the onset of the voltage pulse. Precisely at the resistivity switching from the low-conducting to the high-conducting state at $t = 5.5$ ms the signal becomes constant, which implies that the energy flow is balanced between the electron system, the lattice, and the environment. When the voltage pulse is turned off, the excited charge carriers of the electronic system decay back, and the energy is transferred to the lattice system. The infrared signal remains high, exceeding the duration of the applied voltage, before the lattice system is cooled down linearly in accordance with Newton cooling, as discussed above.

III. HOT ELECTRON MODEL

To explain our observations, we suggest a model of “hot” electrons or nonequilibrium charge carriers [55–57]. It assumes that the electric power is first stored in the electron system causing an increase of the electron temperature T_e . The coupling to the lattice subsystem is responsible for heating up the lattice and increasing the lattice temperature T_L . Two types of carriers are assumed with rather different mobilities. More details are presented in the Supplemental Material [45].

A. Simulation of time dependence

The electron temperature T_e can be calculated by considering energy conservation:

$$nC_e \frac{dT_e}{dt} = P - K, \quad (1)$$

where $P = \sigma(T_e)E_{\text{sample}}^2$ describes the power gained by the electrons, and $K = -\lambda_{\text{therm},e}(T_e - T_L)$ corresponds to the energy transfer from the electrons to the lattice subsystem [58]. We have solved Eq. (1) numerically by first calculating the right side and subsequently determining T_e from the left

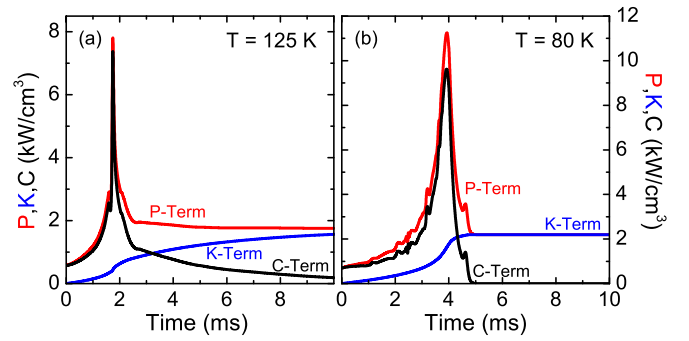


FIG. 12. Time evolution of the input power P (red), the cooling power K (blue), and the power absorbed by the electronic system (black) calculated for the temperature (a) $T = 125 \text{ K}$ and (b) $T = 80 \text{ K}$, where the applied electric field was $E_{\text{sample}} = 250$ and 3000 V/cm , respectively.

side for each time step of the voltage pulse. The electrical conductivity $\sigma(T_e)$ is taken from dc measurements plotted in Fig. S2; the load resistance R_L was experimentally selected between 1 and $100 \text{ k}\Omega$. In addition, we choose an electronic heat capacity $nC_e = 0.2 \text{ J K}^{-1} \text{ cm}^{-3}$ and thermal conductivity of the electron system $\lambda_{\text{therm},e} = 40 \text{ W K}^{-1} \text{ cm}^{-3}$, in agreement with previous investigations, where a similar two-state model was applied to other low-dimensional organic conductors [47,53].

In Fig. 12, the time dependences of the C , P , and K terms are plotted for $T = 125$ and 80 K . The electric power is transferred to the electronic system right from the beginning of the voltage pulse. The cooling term, however, increases only slowly for the elevated temperature $T = 125 \text{ K}$. Before the switching point, the electric power as well as the stored energy in the electric system diverge and drop afterward to a fixed value. The C term decreases slowly to zero since the cooling term approaches the electric power term P initializing the steady state. At low temperatures, $T = 80 \text{ K}$, the power term behaves similarly; only the cooling term K immediately after the resistivity switching reaches the value of the electric input P term. Correspondingly, no energy is then transferred to the electronic system.

The time evolution of the electron temperature T_e is displayed in Fig. 13 for the two experiments at different temperatures. When starting at $T = 125 \text{ K}$, the electron temperature increases more than linearly, then reveals a steplike feature at the phase transition around 2 ms, and finally increases sublinearly up to 160 K at the end of the pulse. For the low-temperature experiment ($T = 80 \text{ K}$), the temperature rises steeply up to $T = 134 \text{ K}$ and stays constant, right below the transition temperature T_{CO} .

The total heat capacity [59] of $\alpha\text{-(BEDT-TTF)}_2\text{I}_3$ crystals below T_{CO} , $C_{\text{tot}} = 0.5 \text{ J K}^{-1} \text{ g}^{-1}$, is more than five times larger than the values for the electron system $C_e = 0.09 \text{ J K}^{-1} \text{ g}^{-1}$ used in our simulations. This agrees well with the fact that phonons typically dominate above 10 K due to the T^3 power law of the lattice contribution compared to the linear dependence of the electronic heat capacity. Assuming that both contributions are independent and add up, $C_{\text{total}} = C_e + C_L \approx 5C_e$, energy conservation yields [53]

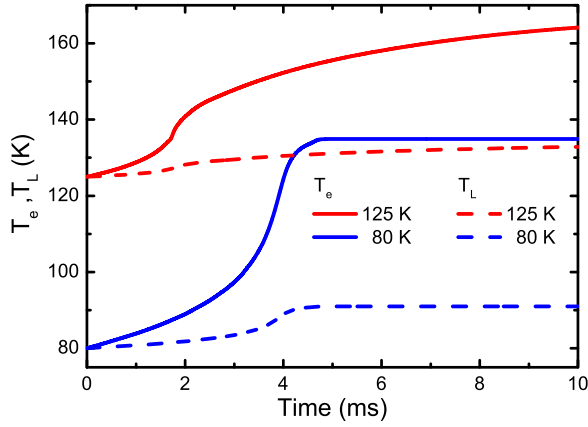


FIG. 13. The electron temperature T_e (solid lines) and the lattice temperature T_L (dashed lines) plotted as a function of time for the two experimental temperatures $T = 125$ K (red) and $T = 80$ K (blue) by applying an external electric field of $E_{\text{sample}} = 250$ and 3000 V/cm, respectively.

$nC_e(T_e - T_L) = nC_L(T_L - T_0)$ and thus

$$\frac{C_e}{C_L} = \frac{(T_L - T_0)}{(T_e - T_L)} = \frac{1}{4} \quad (2)$$

with the initial environmental temperature T_0 ; here T_e is the effective electron temperature and T_L denotes the lattice temperature due to heating.

Using Eq. (2), we estimate the time dependence of the lattice temperature of α -(BEDT-TTF) $_2$ I $_3$ for $T = 125$ and 80 K, and we display the results in Fig. 13. Trailing the electron temperature, the lattice temperature increases only slightly from $T = 125$ to 132 K within the pulse duration of 10 ms; hence it stays below the charge-ordering temperature of $T_{\text{CO}} = 135$ K. Of course, we cannot exclude the idea that due to some scattering centers T_L can locally exceed the transition temperature. We now understand why the reflectivity $\Delta_r R(\nu, t)$ still increases after switching to the high-conducting state, i.e., the lattice temperature T_L keeps rising well after the system has switched. From Fig. 9 we clearly see a metallic transition in the 125 K run, but at $T = 180$ K the reflectivity variation indicates only a small increase of T_L of less than 10 K. This is exactly what our simulation reproduces: the lattice temperature first increases but then saturates as the system switches from the low-conducting to the high-conducting state. This explains why $\Delta_r R(\nu, t)$ remains constant in time until the pulse is switched off.

B. Simulation of field dependence

Up to now, only the time-dependent behavior of the power and the different temperatures were discussed. For a better comparison with our experiments, we have also calculated the current density $J(E, t)$ when the applied electric field is varied. In Fig. 14 the time dependence $J(t)$ is plotted for $T = 125$ and 80 K when the external electric field is varied from 0 to 250 and 3000 V/cm, respectively. Note that J is inversely proportional to the sample resistance R , thus Fig. 14 can be readily compared with the presentation of our experimental transport results in Fig. 3.

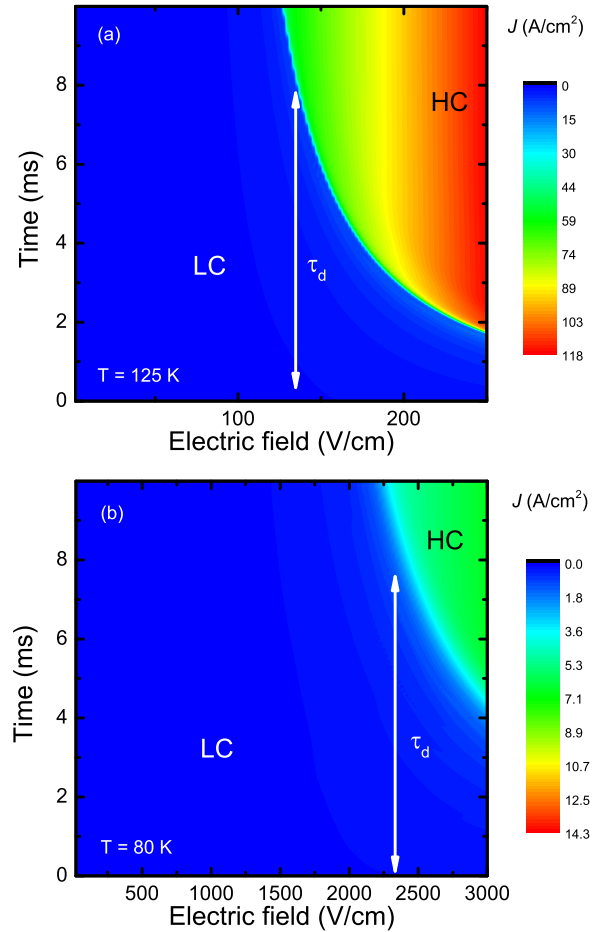


FIG. 14. Variation of the current density $J(E, t)$ with increasing electric field E_{sample} applied to the sample. Simulation for the temperature (a) $T = 125$ K and (b) $T = 80$ K; note the different scales. The arrows indicate the delay time τ_d . The low-conducting state (LC, blue area) is characterized by a small current density, while in the high-conducting part (HC, green-red) the current density is extremely high.

A very distinct jump of $J(E, t)$ occurs when $E_{\text{sample}} > 100$ V/cm providing evidence for the transition from a low-conducting to a high-conducting state; this is in excellent accord with the experimental observations. Also the field-dependent behavior of the delay time τ_d describes the experimental behavior perfectly. In the simulation, the switching occurs at 2000 V/cm $^{-1}$, while our measurements yield $E_{\text{sample}} = 2900$ V/cm. The small deviations between simulations and experiments may be due to the neglect of the contact resistance in our two-point measurement [Fig. 2(a)]; it influences E_{sample} and its T dependence. Furthermore, we assumed a temperature-independent specific heat nC_p and thermal conductivity $\lambda_{\text{therm}, e}$. Nevertheless, we can conclude that the two-state model perfectly reflects our experimental findings and yields an excellent qualitative and quantitative description.

In a next step, the J - E characteristic is calculated to investigate the nonlinear conductivity and the negative differential resistance (NDR). The results are depicted in Fig. 15 for temperatures between $T = 80$ and 130 K. The overall

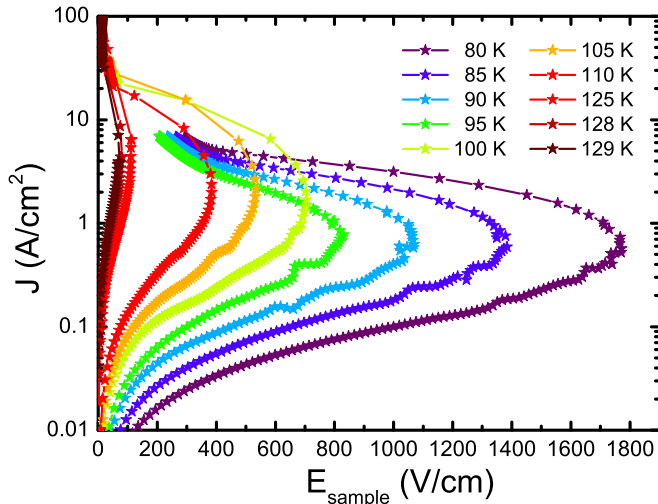


FIG. 15. Simulated J - E characteristic between 0 and 1900 V/cm from temperatures $T = 130$ to 80 K. All curves exhibit a nonlinear regime with a negative differential resistance. Upon cooling, the threshold current decreases while the threshold electric field rises.

behavior is very similar to the data plotted in Fig. 4 resembling an S-shaped current-voltage curve. All temperatures reveal a linear regime at low fields. They exhibit a turning point that marks the threshold field E_{thresh} and current density J_{thresh} . All J - E curves cross over in the NDR regime. Above a certain current density, the field increases again but much steeper than for lower electric fields, indicating that the high-field conduction state is different from the initial state at low fields. Small deviations from the experimental findings can be noted, such as slightly higher current density after the NDR compared to the experiment, or the higher fields for which the curves turn. We attribute these to the unknown contact resistance in the experiment, which is connected in series to the sample and load resistance and affects E_{sample} and J directly. Nevertheless, the detailed agreement of the simulation with the experiment is remarkable.

Following our analysis of the experimental data in Sec. II A, we have extracted the threshold current J_{thresh} and the threshold field E_{thresh} from the J - E curves in Fig. 15 and plotted them as a function of temperature in Fig. 16. The results of our simulations resemble more or less closely the experimental data displayed in Fig. 5. The observed threshold field $E_{\text{thresh}}(T)$ was well described by a linear increase as the temperature is reduced below T_{CO} . Our simulation, however, rises in a quadratic manner. This may be attributed to the decrease of the occupation number of the conduction band upon cooling. Moreover, the increase of E_{thresh} at low temperatures can also be attributed to thermal depletion of the conduction band upon cooling.

The calculated current density, on the other hand, decays linearly upon cooling, while the experiments seem to flatten at low temperatures. However, the threshold power density $P_{\text{thresh}}(T)$ increases steeply close to the phase transition and crosses over in the regime with a smaller slope, which resembles the development of the energy gap.

We have seen that the two-state model excellently describes the electrically induced phase transition and explains qualitatively and, impressively, also the quantitative behavior of the

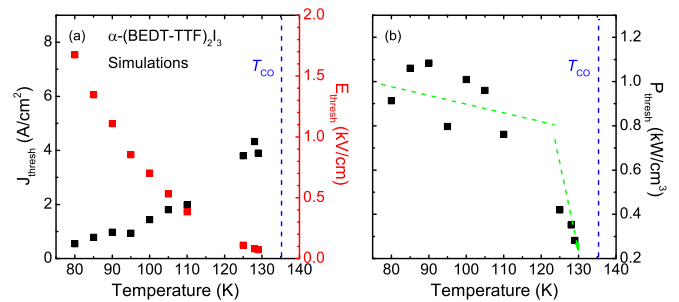


FIG. 16. (a) Numerical simulations of the threshold electric field (red) and current density (black) as a function of temperature below T_{CO} using parameters for α -(BEDT-TTF) $_2$ I $_3$. (b) Temperature dependence of the threshold power P_{thresh} determined from the product of E_{thresh} and J_{thresh} . The vertical dotted line marks the phase-transition temperature of $T_{\text{CO}} = 135$ K.

experimental values. It not only supports the time-resolved transport measurements, but it also explains in detail the observations made in the time-dependent reflectivity study. It confirms that we are dealing with an electronic system linearly coupled to the crystal lattice. Furthermore, we made a precise estimation of the lattice temperature T_L that is different from the electronic system and agrees nicely with the estimated sample temperature derived from the comparison of time-dependent reflectivity with the steady-state spectra.

C. Field-induced Dirac-like electrons

We have demonstrated that switching to a high-conducting state is caused by excitations of charge carriers into a state with a high mobility. Here, the question arises whether the enhanced mobility is somehow related to the linear band dispersion found in α -(BEDT-TTF) $_2$ I $_3$. The tilted Dirac cone, i.e., the touching of the bands at the Fermi energy, is only predicted for high pressure. Nevertheless, band-structure calculations (Fig. S4) yield that the linear dispersion exists also in the insulating phase [45,60]. Furthermore, the temperature dependence of the resistance under pressure normalized to the room-temperature values exhibits exactly the same slope as the resistance at ambient pressure [22,61]. Since Dirac fermions are made responsible for the conductivity behavior under pressure, we propose that they also contribute to the metallic state at ambient conditions, excited by a strong electric field. The coexistence of two different types of charge carriers with two orders of magnitude different mobility was previously suggested based on magnetotransport [62] and pressure-dependent optical experiments [27] in α -(BEDT-TTF) $_2$ I $_3$. We propose that also the electrons excited in the conduction band can act either as normal massive electrons or as massless Dirac-like electrons with very high mobility. This picture provides a consistent explanation of the results presented here.

IV. CONCLUSIONS

We have reported comprehensive time-resolved investigations of the electrodynamics at the metal-insulator transition in α -(BEDT-TTF) $_2$ I $_3$. In the charge-ordered state below $T_{\text{CO}} = 135$ K, the time- and field-dependent charge transport was measured by applying a voltage pulse along the a direction.

We found an electrically induced resistivity switching from a low-conducting to a high-conducting state, leading to an S-like shape in the J - E curve. In a regime of negative differential resistance, the electric field across the sample drops drastically above a certain threshold, which increases upon cooling.

Measuring the polarization-dependent infrared reflectivity after a strong electric field was applied allows us to study the transient optical properties in a wide temperature and field range. From the spectral signature at $T = 125$ K, we can identify a field-induced metallic state similar to the high-temperature conducting phase. In contrast, at temperatures well into the insulating state ($T = 80$ K), the reflectivity variations are completely different and cannot be explained by Joule heating. We suggest the creation of an electronically driven high-conducting state in α -(BEDT-TTF)₂I₃. These field-excited charge carriers exhibit an extremely high mobility known from the Dirac-like electrons present under high pressure. Thus we propose that the electrons excited in the conduction band can act either as normal massive electrons or as massless Dirac-like electrons with very high mobility.

This generation of hot charge carriers by the electric field is then simulated with a two-state model. Only based on energy conservation and known material parameters, our numerical simulations can—qualitatively as well as quantitatively—reproduce the experimentally observed properties, such as the temporal behavior of the resistivity as a function of the electric field, the negative differential resistivity, and the different threshold values. By defining an effective electron temperature T_e , we estimate the lattice temperature T_L close to the measured sample temperature. For all parameters analyzed, the quantitative agreement between model and experiment is excellent.

ACKNOWLEDGMENTS

We would like to thank R. Beyer, E. Rose, D. Wu, and S. Zapf for useful discussions, as well as G. Untereiner for technical support. Funding by the Deutsche Forschungsgemeinschaft (DFG) and Deutscher Akademischer Austauschdienst (DAAD) is acknowledged.

-
- [1] K. Bender, H. Endres, H.-W. Helberg, I. Hennig, H. Schäfer, H. Keller, and D. Schweitzer, *Mol. Cryst. Liq. Cryst.* **107**, 45 (1984).
 - [2] M. Dressel, G. Grüner, J. Pouget, A. Breining, and D. Schweitzer, *J. Phys. (Paris) I* **4**, 579 (1994).
 - [3] H. Kino and H. Fukuyama, *J. Phys. Soc. Jpn.* **64**, 1877 (1995).
 - [4] H. Kino and H. Fukuyama, *J. Phys. Soc. Jpn.* **65**, 2158 (1996).
 - [5] H. Seo, C. Hotta, and H. Fukuyama, *Chem. Rev.* **104**, 5005 (2004).
 - [6] Y. Takano, K. Hiraki, H. Yamamoto, T. Nakamura, and T. Takahashi, *J. Phys. Chem. Solids* **62**, 393 (2001).
 - [7] T. Takahashi, Y. Nogami, and K. Yakushi, *J. Phys. Soc. Jpn.* **75**, 051008 (2006).
 - [8] J. Moldenhauer, C. Horn, K. Pokhodnia, D. Schweitzer, I. Heinen, and H. Keller, *Synth. Met.* **60**, 31 (1993).
 - [9] R. Wojciechowski, K. Yamamoto, K. Yakushi, M. Inokuchi, and A. Kawamoto, *Phys. Rev. B* **67**, 224105 (2003).
 - [10] M. Dressel and N. Drichko, *Chem. Rev.* **104**, 5689 (2004).
 - [11] Y. Yue, K. Yamamoto, M. Uruichi, C. Nakano, K. Yakushi, S. Yamada, T. Hiejima, and A. Kawamoto, *Phys. Rev. B* **82**, 075134 (2010).
 - [12] T. Ivek, B. Korin-Hamzić, O. Milat, S. Tomić, C. Clauss, N. Drichko, D. Schweitzer, and M. Dressel, *Phys. Rev. B* **83**, 165128 (2011).
 - [13] K. Yakushi, *Crystals* **2**, 1291 (2012).
 - [14] T. Kakiuchi, Y. Wakabayashi, S. Hiroshi, T. Takahashi, and T. Nakamura, *J. Phys. Soc. Jpn.* **76**, 113702 (2007).
 - [15] T. Ivek, B. Korin-Hamzić, O. Milat, S. Tomić, C. Clauss, N. Drichko, D. Schweitzer, and M. Dressel, *Phys. Rev. Lett.* **104**, 206406 (2010).
 - [16] S. Tomić and M. Dressel, *Rep. Progr. Phys.* **78**, 096501 (2015).
 - [17] P. Lunkenheimer, B. Hartmann, M. Lang, J. Müller, D. Schweitzer, S. Krohns, and A. Loidl, *Phys. Rev. B* **91**, 245132 (2015).
 - [18] N. Tajima, A. Ebina-Tajima, M. Tamura, Y. Nishio, and K. Kajita, *J. Phys. Soc. Jpn.* **71**, 1832 (2002).
 - [19] A. Kobayashi, S. Katayama, K. Noguchi, and Y. Suzumura, *J. Phys. Soc. Jpn.* **73**, 3135 (2004).
 - [20] M. Dressel, G. Grüner, J. Pouget, A. Breining, and D. Schweitzer, *Synth. Met.* **70**, 929 (1995).
 - [21] T. Ivek, I. Kovačević, M. Pinterić, B. Korin-Hamzić, S. Tomić, T. Knoblauch, D. Schweitzer, and M. Dressel, *Phys. Rev. B* **86**, 245125 (2012).
 - [22] N. Tajima, S. Sugawara, M. Tamura, Y. Nishio, and K. Kajita, *J. Phys. Soc. Jpn.* **75**, 051010 (2006).
 - [23] T. Mori, *J. Phys. Soc. Jpn.* **79**, 014703 (2010).
 - [24] N. Tajima, S. Sugawara, M. Tamura, R. Kato, Y. Nishio, and K. Kajita, *Europhys. Lett.* **80**, 47002 (2007).
 - [25] Y. Suzumura and A. Kobayashi, *Crystals* **2**, 266 (2012).
 - [26] K. Kajita, Y. Nishio, N. Tajima, Y. Suzumura, and A. Kobayashi, *J. Phys. Soc. Jpn.* **83**, 072002 (2014).
 - [27] R. Beyer, A. Dengl, T. Peterseim, S. Wackerow, T. Ivek, A. V. Pronin, D. Schweitzer, and M. Dressel, *Phys. Rev. B* **93**, 195116 (2016).
 - [28] N. Tajima, J. Fujisawa, N. Naka, T. Ishihara, R. Kato, Y. Nishio, and K. Kajita, *J. Phys. Soc. Jpn.* **74**, 511 (2005).
 - [29] S. Iwai, K. Yamamoto, A. Kashiwazaki, F. Hiramatsu, H. Nakaya, Y. Kawakami, K. Yakushi, H. Okamoto, H. Mori, and Y. Nishio, *Phys. Rev. Lett.* **98**, 097402 (2007).
 - [30] Y. Kawakami, T. Fukatsu, Y. Sakurai, H. Unno, H. Itoh, S. Iwai, T. Sasaki, K. Yamamoto, K. Yakushi, and K. Yonemitsu, *Phys. Rev. Lett.* **105**, 246402 (2010).
 - [31] S. Iwai, *Crystals* **2**, 590 (2012).
 - [32] K. Yamamoto, S. Iwai, S. Boyko, A. Kashiwazaki, F. Hiramatsu, C. Okabe, N. Nishi, and K. Yakushi, *J. Phys. Soc. Jpn.* **77**, 074709 (2008).
 - [33] K. Yamamoto, A. Kowalska, and K. Yakushi, *Appl. Phys. Lett.* **96**, 122901 (2010).
 - [34] K. Yamamoto, A. A. Kowalska, and K. Yakushi, *Phys. Status Solidi C* **9**, 1189 (2012).
 - [35] T. Naito, *Molecular Electronic and Related Materials—Control and Probe with Light* (Transworld Research Network, Kerala, India, 2010).

- [36] T. Iimori and N. Ohta, *J. Phys. Chem. C* **118**, 7251 (2014).
- [37] H. Nakaya, F. Hiramatsu, Y. Kawakami, S. Iwai, K. Yamamoto, and K. Yakushi, *J. Lumin.* **128**, 1065 (2008).
- [38] T. Iimori, T. Naito, and N. Ohta, *J. Phys. Chem. C* **113**, 4654 (2009).
- [39] T. Iimori, T. Naito, and N. Ohta, *Chem. Lett.* **36**, 536 (2007).
- [40] T. Iimori, T. Naito, and N. Ohta, *J. Am. Chem. Soc.* **129**, 3486 (2007).
- [41] K. Tamura, T. Ozawa, Y. Bando, T. Kawamoto, and T. Mori, *J. Appl. Phys.* **107**, 103716 (2010).
- [42] S. Katayama, A. Kobayashi, and Y. Suzumura, *J. Phys. Soc. Jpn.* **75**, 054705 (2006).
- [43] P. Alemany, J.-P. Pouget, and E. Canadell, *Phys. Rev. B* **85**, 195118 (2012).
- [44] K. Bender, I. Hennig, D. Schweitzer, K. Dietz, H. Endres, and H. Keller, *Mol. Cryst. Liq. Cryst.* **108**, 359 (1984).
- [45] See Supplemental Material at <http://link.aps.org/supplemental/10.1103/PhysRevB.93.245133> for details on the experimental setup, a description of the transport measurements, estimates of the Joule heating, and a description of the band structure and the two-state model.
- [46] F. Itose, T. Kawamoto, and T. Mori, *J. Appl. Phys.* **113**, 213702 (2013).
- [47] T. Ozawa, K. Tamura, Y. Bando, T. Kawamoto, T. Mori, and I. Terasaki, *Phys. Rev. B* **80**, 155106 (2009).
- [48] G. Grüner, *Rev. Mod. Phys.* **60**, 1129 (1988).
- [49] K. Maki, *Phys. Rev. B* **33**, 2852 (1986).
- [50] M. Dressel and G. Grüner, *Electrodynamics of Solids* (Cambridge University Press, Cambridge, 2002).
- [51] C. Clauss, N. Drichko, D. Schweitzer, and M. Dressel, *Physica B* **405**, S144 (2010).
- [52] T. Peterseim and M. Dressel, *J. Infrared Millim. Terahertz Waves*, doi:10.1007/s10762-016-0294-5 (to be published) [arXiv:1604.00977].
- [53] T. Mori, T. Ozawa, Y. Bando, T. Kawamoto, S. Niizeki, H. Mori, and I. Terasaki, *Phys. Rev. B* **79**, 115108 (2009).
- [54] A. Zimmers, L. Aigouy, M. Mortier, A. Sharoni, S. Wang, K. G. West, J. G. Ramirez, and I. K. Schuller, *Phys. Rev. Lett.* **110**, 056601 (2013).
- [55] D. M. Kroll, *Phys. Rev. B* **9**, 1669 (1974).
- [56] S. M. Sze and K. K. Ng, *Physics of Semiconductor Devices*, 3rd ed. (Wiley-Interscience, Hoboken, NJ, 2007).
- [57] P. Y. Yu and M. Cardona, *Fundamentals of Semiconductors: Physics and Materials Properties*, 4th ed. (Springer, Berlin, 2010).
- [58] For our calculations, we set the lattice temperature T_L equal to the environment temperature T_0 . The last term corresponds to the experimental situation described by Figs. 11(b) and 11(d): the reflectivity does not change, and correspondingly the sample has a constant temperature. We have assumed a linear Newton cooling.
- [59] N. A. Fortune, K. Murata, M. Ishibashi, M. Tokumoto, N. Kinoshita, and H. Anzai, *Solid State Commun.* **79**, 265 (1991).
- [60] T. Peterseim and M. Dressel, arXiv:1603.07505.
- [61] H. Schwenk, F. Gross, C.-P. Heidmann, K. Andres, D. Schweitzer, and H. Keller, *Mol. Cryst. Liq. Cryst.* **119**, 329 (1985).
- [62] M. Monteverde, M. O. Goerbig, P. Auban-Senzier, F. Navarin, H. Henck, C. R. Pasquier, C. Mézière, and P. Batail, *Phys. Rev. B* **87**, 245110 (2013).