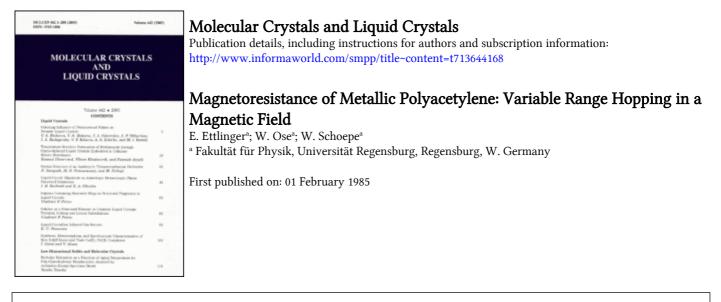
This article was downloaded by: *[German National Licence 2007]* On: *20 December 2010* Access details: *Access Details: [subscription number 777306419]* Publisher *Taylor & Francis* Informa Ltd Registered in England and Wales Registered Number: 1072954 Registered office: Mortimer House, 37-41 Mortimer Street, London W1T 3JH, UK



**To cite this Article** Ettlinger, E., Ose, W. and Schoepe, W.(1985) 'Magnetoresistance of Metallic Polyacetylene: Variable Range Hopping in a Magnetic Field', Molecular Crystals and Liquid Crystals, 117: 1, 173 – 176 **To link to this Article: DOI:** 10.1080/00268948508074618

URL: http://dx.doi.org/10.1080/00268948508074618

## PLEASE SCROLL DOWN FOR ARTICLE

Full terms and conditions of use: http://www.informaworld.com/terms-and-conditions-of-access.pdf

This article may be used for research, teaching and private study purposes. Any substantial or systematic reproduction, re-distribution, re-selling, loan or sub-licensing, systematic supply or distribution in any form to anyone is expressly forbidden.

The publisher does not give any warranty express or implied or make any representation that the contents will be complete or accurate or up to date. The accuracy of any instructions, formulae and drug doses should be independently verified with primary sources. The publisher shall not be liable for any loss, actions, claims, proceedings, demand or costs or damages whatsoever or howsoever caused arising directly or indirectly in connection with or arising out of the use of this material.

Mol. Cryst. Liq. Cryst. 1985, Vol. 117, pp. 173-176 0026-8941/85/1174-0173/\$10.00/0 © 1985 Gordon and Breach, Science Publishers, Inc. and OPA Ltd. Printed in the United States of America

> MAGNETORESISTANCE OF METALLIC POLYACETYLENE:VARIABLE RANGE HOPPING IN A MAGNETIC FIELD

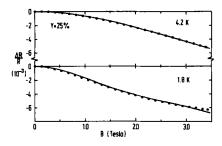
E. ETTLINGER, W. OSE, W. SCHOEPE Fakultät für Physik, Universität Regensburg D-8400 Regensburg, W.Germany

Abstract The magnetoresistance (MR) of heavily doped (CH) x has been measured between 4.2 K and 0.3 K in fields up to 3.4 Tesla. The data can be described by variable-range hopping (VRH) in a magnetic field and by postulating a second transport mechanism independent of the magnetic field. A model based on orbital shrinking of the localized wavefunction (positive MR) and on a Zeeman shift of the energy levels (negative MR) can be fit to the data and yields the Bohr radius and the binding energy of the localized states as well as the variation of the density of states at the Fermi level, which in most samples is found to be quadratic.

We have studied the dc conductivity of heavily doped polyacetylene  $(I_3, AsF_6)$  at temperatures between 4.2 K and 0.3 K and in magnetic fields up to 3.4 Tesla. From the MR in particular it is possible to obtain detailed information on the transport mechanism.

Our samples were polymerized by the Stuttgart group. Iodine doping was performed in Regensburg by exposing the samples to iodine vapor and consecutive vacuum pumping. The magnetic field was oriented parallel to the polyacetylene films and to the current.

Typical results are shown in Figs. 1 and 2 where the relative change of the resistance as a function of the magnetic field is displayed at various constant temperatures. At high temperatures (Fig. 1) the MR is negative but upon cool-down continuously changes to positive values (Fig. 2). At the lowest temperatures the positive MR is similar to our earlier data on less metallic samples<sup>1</sup>, the differences being the two orders of magnitude smaller values



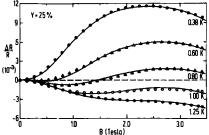


Fig. 1 Negative MR of  $(CHI_{0.25})_x$  at high temperatures. The solid x lines are fits of Eq.(1).

Fig. 2 Change from negative to positive MR at lower temperature (same sample as in Fig. 1). The solid lines are fits of Eq.(1).

and the maximum near 2.5 Tesla instead of a monotonic increase.

In accordance with the interpretation of our previous work we attribute the positive part of the MR to orbital shrinking<sup>2</sup> of the hole wavefunction resulting in a reduction of the conductivity by the factor  $\exp((B/B_0)^2$  in moderate fields or  $\exp((B/B_1)^{1/4}$  in high fields<sup>3</sup>. The characteristic fields  $B_0$  and  $B_1$  have particular temperature dependencies, which are determined by the variation of the density of states N at the Fermi level  $E_F$  (see Table I).

For intermediate fields we use an empirical interpolation formula  $\exp(-f(B))$  where  $f(B) = \frac{B^2}{(B_o^2 + B_1^{1/4} B^{7/4})}$ . Hence the VRH conductivity in a magnetic field is given by

 $\sigma_{\rm VRH}(B) = \sigma_{\rm VRH}(0) \cdot \exp(-f(B)).$ 

The negative part of the MR can be described by a Zeeman shift of the energy levels of the hole wavefunction, which leads to a spin dependent Bohr radius<sup>4</sup>. The overall effect is an increase of the conductivity. In the limit of a Zeeman shift being much smaller than the binding energy one finds<sup>4</sup> the factor  $ch(B/B_2)$ , where the temperature dependence of the characteristic field  $B_2$  again depends upon the density of states (see Table I). Taking into account both contributions we write

$$\sigma_{VRH}(B) = \sigma_{VRH}(0) \cdot \exp(-f(B)) \cdot ch(B/B_2).$$

In addition, we postulate a second transport mechanism  $\sigma_2$  being insensitive to a magnetic field, which in the limit of low temperatures and high fields shortens out the vanishing VRH and thus accounts for the saturation tendency of our data in this regime:  $\sigma(B) = \sigma_{VRH}(B) + \sigma_2$ . From the total conductivity we finally obtain for the MR

$$\frac{\Delta R}{R} = c \cdot (1 - \exp(-f(B)) \cdot ch(B/B_2)), \qquad (1)$$

with c =  $\sigma_{VRH}(0)/(\sigma_{VRH}(0) + \sigma_2)$ .

The solid lines in Figs. 1 and 2 are fits of equation (1) to the data. Fitting parameters are c, the temperature independent coefficients of the characteristic fields  $B_i$  (i=0,1,2), and the power law which determines their temperature dependence.

The results for two of our samples  $(8\% I_3 = 11\% AsF_6)$  are compiled in Table I together with a listing of theoretical predictions for purely constant and quadratic density of states N.

Table I	Temperature	dependence	of	the	characteristic	fields
---------	-------------	------------	----	-----	----------------	--------

Characteristic			Experiment		
fields	N = const.	$N=N_o(E-E_F)^2$	AsF <sub>6</sub>	1 <u>3</u>	
В	<sub>T</sub> 3/8	<sub>7</sub> 3/4	_0.47	т <sup>0.65</sup>	
о В,	$\frac{1}{T}^{2/3}$	$r^2$		_1.73	
<sup>2</sup> 1 В <sub>2</sub>	$\frac{1}{r^{1/4}}$	$T^{1/2}$			

For the iodine doped sample we find the following values at 1 K:  $B_0 = 1.1$  Tesla,  $B_1 = 1.0 \cdot 10^{-3}$  Tesla, and  $B_2 = 0.69$  Tesla.

A comparison between theory and experiment in Table I shows that for the iodine doped sample the temperature dependences of the  $B_i$  (i = 0,1,2) indicate a parabolic density of states at the Fermi level as observed earlier<sup>1</sup> in contrast to the  $AsF_6$  doped sample, where N seems to be nearly constant. At present it is unclear whether the differences in the density of states are caused by the different dopants or by some other properties of the various samples.

From the coefficients of the three characteristic fields one can in principle derive the following three microscopic quantities: 1. The Bohr radius a of the hole, 2. it's binding energy  $\Delta E$ , and 3. the factor N<sub>o</sub> of the density of states. This, however, requires the knowledge of numerical factors which theories of VRH leave uncertain to within an order of magnitude. Therefore, we obtain only estimates as follows (for the iodine doped sample):  $a \approx 100$  Å,  $\Delta E \approx 2$  K, N<sub>o</sub>  $\approx 10^{26}$  (eV cm)<sup>-3</sup>. These values do not appear to be unreasonable.

In summary, the MR of heavily doped polyacetylene can be described by 3D variable-range hopping of localized states with spin. Further work is in progress to determine the influence of the dopant concentration on the properties of the hole. Even more interesting would be an understanding of the nature of the  $\sigma_2$  conduction mechanism.

We should like to thank the Stuttgart group, especially K. Ehinger and S. Roth, for supplying the samples. E.E. is supported by a Bavarian Graduate Scholarship and W.S. by the Deutsche Forschungsgemeinschaft through a Heisenberg grant.

## References

- E. Ettlinger, W. Schoepe, M. Monkenbusch, and G. Wieners, Solid State Commun. 49, 107 (1984).
- For a review see B.I. Shklovskii, Fiz.Tekh.Poluprov. <u>6</u>, 1197 (1972) [Sov.Phys. Semicond. <u>6</u>, 1053 (1973)].
- B.I. Shklovskii, Fiz.Tekh.Poluprov. <u>17</u>, 2055 (1983) [(Sov.Phys.Semicond. <u>17</u>, 1311 (1983)]. With a constant density of states we get for the argument -(B/B<sub>1</sub>)<sup>3/8</sup>, where B<sub>1</sub> ∝ T<sup>2/3</sup>.
  H. Fukuyama and K. Yosida, J.Phys.Soc. Japan <u>46</u>, 102 (1979);
- H. Fukuyama and K. Yosida, J.Phys.Soc. Japan <u>46</u>, 102 (1979); <u>46</u>, 1522 (1979).