# MAGNETIC SUSCEPTIBILITY OF PbTe AND PbSe\*\*\*\*

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(Received May 20, 1988)

The magnetic susceptibility of Bridgman-grown single crystals of PbTe and PbSe has been examined. The measurements were carried out using a SQUID detection system over a temperature range from 2 to 100 K in magnetic fields up to 1 T. Both systems were diamagnetic at 100 K with a susceptibility of about  $-3.6 \times 10^{-7}$  emu/g for PbSe and -3.1x10<sup>-7</sup> emu/g for PbTe. With decreasing temperature the magnitude of the susceptibility increased slightly; below 10 K there was a rapid decrease, an order of magnitude for PbTe and about 10% for PbSe.

PACS numbers: 75.20.Ck

# 1. Introduction

Recently, we have measured magnetic properties of IV-VI diluted magnetic semiconductors, such as  $Pb_{1-x}Mn_xTe$ ,  $Pb_{1-x}Gd_xTe$ , and  $Pb_{1-x}Mn_xSe$  [1-3]. In order to such as  $Pb_{1-x}Mn_xTe$ ,  $Pb_{1-x}Gd_xTe$ , and  $rb_{1-x}Mn_xee$  it was necessary to know the magnetization and susceptibility measurements it was necessary to know the the magnetization and susceptibility measurements it was necessary to magnetic contribution of the host material. Therefore, we have measured the magnetic susceptibility of the host material. to 1 T of PbTe from 1.8 K to 250 K and of PbSe from 2.2 K to 100 K at fields from

Proc. XVII School on Physics of Semiconducting Compounds, Jaszowiec 1988.

This word the U.S. ARO under

This work was supported in part by the U.S. DARPA and the U.S. ARO under Grant No. 29.85-K-0052

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The PbTe and PbSe samples were cut out from larger boules grown by the Bridge technique. Transport parameters at liquid helium temperature are shown in Table 1977.

We had two different types of PbTe samples. In samples of PbTe 2 the carrier centration, determined by assuming one-carrier transport, had an unrealistic, low at 4.2 K. This, together with the low mobility value, suggested a strong compensation unknown defects. The samples of PbTe 1 and PbSe were typical pure lead chalcographs.

The susceptibility of single crystals of PbTe and PbSe was measured using a squeetection system. Our experimental set-up is described elsewhere [1–3]. The measurement

TAB

Sample	Туре	Carrier concentration [cm <sup>-3</sup> ]	Hall mobility [cm²/V·s]
PbTe 1	p	4.78×10 <sup>18</sup>	1.01×10 <sup>5</sup>
PbTe 2	n	4.69 × 10 <sup>15</sup>	$5.60 \times 10^{3}$
PbSe	n	$3.06 \times 10^{18}$	$1.29 \times 10^{5}$

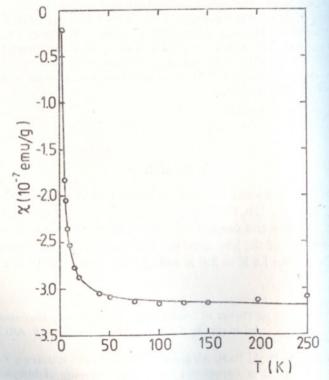


Fig. 1. Magnetic susceptibility vs temperature for PbTe 1. Solid line is a fit of the Curie la

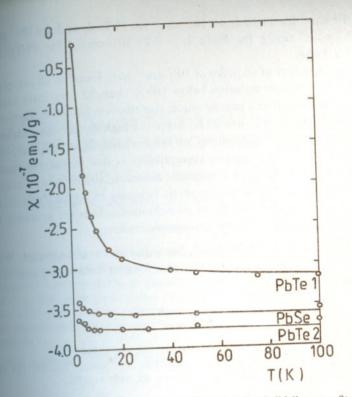


Fig. 2. Magnetic susceptibility vs temperature for PbTe and PbSe. Solid lines are fits of the Curie law

were made on samples of PbTe 1 from 1.8 K to 250 K, of PbTe 2 from 2 K to 100 K, and of PbSe from 2.2 K to 100 K. The magnetization was measured at each temperature at three fields, usually 0.1, 0.5, and 1 T. The magnetic susceptibility was determined by a linear least squares fit with an accuracy of about 2%.

The magnetic susceptibility vs temperature for samples PbTe 1 is shown in Fig. 1, and for PbTe 1, PbTe 2, and PbSe in Fig. 2. All materials were diamagnetic, but there was a strong evidence of paramagnetic contribution below 20 K.

## 3. Discussion

The susceptibility data have been fitted to the Curie law:

$$\chi = \frac{N_{\rm S} g^2 \mu_{\rm B}^2 S(S+1)}{3k_{\rm B} T} + \chi_{\rm dia},\tag{1}$$

where  $N_s$  is a number of spins, S=1/2 is the assumed spin, g=2 is the assumed spin-splitting factor,  $\mu_B$  is the Bohr magneton,  $k_B$  is the Boltzmann constant, T is the absolute temperature and  $\chi_{\rm dia}$  is the lattice diamagnetic susceptibility, assumed to be constant. The results of fits are shown in Figs 1 and 2 as solid lines. The fitting parameters were:

 $N_{\rm S}$  about  $8 \times 10^{18}$  cm<sup>-3</sup> for PbTe 1,  $4 \times 10^{17}$  cm<sup>-3</sup> for PbTe 2, and  $6 \times 10^{17}$  cm<sup>-3</sup> for PbSe;  $\chi_{\rm dia}$  about  $-3.2 \times 10^{-7}$  emu/g for PbTe 1,  $-3.8 \times 10^{-7}$  emu/g for PbTe 2, and -3.65 ×  $10^{-7}$  emu/g for PbSe.

No magnetic impurities of an order of 10<sup>17</sup> cm<sup>-3</sup> were found in all samples by an X-tay fluorescence. The carrier concentration below 100 K changed very little with temperature and it does not seem likely that a paramagnetic contribution from free carriers can account for the paramagnetic behavior below 20 K. Since the high carrier concentration in PbTe1 is probably due to the nonstoichiometry of the material, it seems likely that the paramagnetic contribution to the magnetic susceptibility is due to the presence of Te<sup>-</sup> ions or Pb vacancies which would carry a magnetic moment. Similarly, an excess of Pb<sup>+</sup> ions or Te vacancies would lead to the paramagnetic behavior in n-type PbSe. Samples PbTe 2 seemed to be strongly compensated, the nonstoichiometric ions are probably either neutral or doubly ionized, and do not carry a magnetic moment.

We wish to thank Drs. P. Becla and A. Szczerbakow for the samples. We are grateful for the use of the SQUID facilities at the National Magnet Laboratory and to Dr. R. Frankel for his help.

#### REFERENCES

- [1] J. R. Anderson, M. Górska, Solid State Commun. 52, 601 (1984).
- [2] M. Górska, J. R. Anderson, Z. Gołacki, Mat. Res. Soc. Symp. Proc. 89, 119 (1987).
- [3] M. Górska, J. R. Anderson, Solid State Commun. 63, 1055 (1987).