MS25-P4  
**Thermal and magnetic anomalies of Mn$_{1-x}$Co$_x$Ge**

Gleb A. Valkovskiy$^1$, Evgeniy V. Altyndaev$^{1,2}$, Maria D. Kuchugura$^{1,2}$, Ekaterina G. Yashina$^{1,2}$, Vadim A. Dyadkin$^1$, Anatoly V. Tsvyashchenko$^3$, Dmitry Chernyshov$^4$, Sergey V. Grigoriev$^{1,2}$

1. Faculty of Physics, Saint Petersburg State University, 198504 Saint Petersburg, Russia
2. Petersburg Nuclear Physics Institute, Gatchina, 188300 Saint Petersburg, Russia
3. Swiss-Norwegian Beamlines at the ESRF, Grenoble 38000, France
4. Institute for High Pressure Physics, 142190, Troitsk, Moscow Region, Russia

email: Valkovskyy.Gleb@mail.ru

The helimagnets with chiral spin structure are interesting because of a wide range of unusual phenomena related to their magnetic ordering. An incomplete but representative list includes skyrmion structures, magnetostriction and magnetoresistance, coupling of structural and magnetic chiralities [1-5]; some of them may found application in the next generation spintronic devices. Besides well-known MnSi, this family includes MnGe, possessing a number of advantages, such as high transition temperature $T_N$, of about 170 K (instead of $\approx$ 29 K in MnSi) and a large ordered Mn moment of about 1.8 $\mu_B$ at 2 K (compared to 0.4 $\mu_B$ at 2 K in MnSi) [6 - 8]. It was shown that MnGe has additional specific feature, namely a broad transition region from ordered helical phase to a disordered paramagnetic one, from $T_{N}$ to $T_{N}$ $\approx$ 270 K [6 - 8]. The nature of the region is not yet entirely clear, in particular ferromagnetic nanoregions ($\sim$ 1 nm) were suggested to exist between $T_{N}$ and $T_{N}$, based on small angle neutron scattering data [6]. Here we report an anomaly in thermal expansion in the same region for MnGe and follow the evolution of thermal properties as a function of $x$ in the series of Mn$_x$Co$_{1-x}$Ge solid solutions, i.e. from the helimagnet MnGe to the diamagnet CoGe. Thermal expansion was studied by fitting the lattice parameter dependence on temperature based on synchrotron powder diffraction data in terms of the Debye model. The deviation from the Debye model increases with decreasing cobalt concentration, with the largest deviation for MnGe in the region between $T_N$ and $T_C$. In particular, a negative increment to the linear thermal expansion coefficient was observed for MnGe in this temperature range. The reason for this effect is considered to be anomalous magnetic behavior.

**Keywords:** magnetostriction, powder diffraction

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**Magnetic-crystallographic $p,T$-phase diagram of Fe$_{1-x}$Te: A high-pressure neutron diffraction study**

Jens-Erik Jørgensen$^1$

1. Dept. of Chemistry, Aarhus University, DK-8000 Aarhus C, Denmark

email: jenserik@chem.au.dk

The crystal and magnetic structures of Fe$_{1-x}$Te for $x = 0.087$ and 0.141 have been studied by neutron powder diffraction in the temperature range from 5 to 170 K at pressures in the range from $\approx$0.8 to $\approx$7 GPa. The $p,T$-phase diagrams of the two Fe$_{1-x}$Te compounds contain three phases with monoclinic, orthorhombic and tetragonal symmetry. The tetragonal phase with space group $P4_{2}mn$ is stable at ambient conditions as well as at pressures. In the case of Fe$_{1.087}$Te the monoclinic and orthorhombic phases are both antiferromagnetically ordered and stable at temperatures below $\approx$69 K while the non-magnetic tetragonal phase is stable above this temperature. The monoclinic phase is stable for $p < 1.2$ GPa while the orthorhombic phase is stable for $1.2 < p < 1.7$ GPa and the tetragonal phase becomes stable at higher pressures at the lowest measured temperatures. The $p,T$-phase diagrams of Fe$_{1-x}$Te are shown in Fig. 1. Fe$_{1.141}$Te shows the same type of $p,T$-phase diagram as Fe$_{1.087}$Te although the stability range of the orthorhombic is larger, $0.8 \leq p \leq 2.3$ GPa. The magnetic ordering is antiferromagnetic bicollinear and commensurate with propagation vector $\mathbf{k} = (1/2 \ 0 \ 1/2)$ in the monoclinic phase while it is incommensurate with propagation vector $\mathbf{k} = (1/2 - 0 \ 1/2)$ in the orthorhombic phase. The wave-length of the modulation of the magnetic structure of the orthorhombic phases was found to increase with pressure. The pressure-induced collapse of magnetic order at $\approx$1.7 GPa in the case of Fe$_{1.087}$Te is accompanied by an abrupt change in volume and compressibility, suggestive of a spin state change of the Fe$^{2+}$ ions in the FeTe layers. The observed abrupt change in volume and compressibility is presumably related to the earlier observed transition from the tetragonal to a collapsed tetragonal phase at $\approx$4 GPa at ambient temperature [1].