

Metastability of crystal and magnetic structures in compounds containing heavy rare earths

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A comprehensive review is given of our recent advances in the study of metastable phases in intermetallics RCu_5 and $\text{RGe}_6\text{Fe}_{6-x}\text{Mn}_x$ ($x=0, 3, 6$) containing heavy rare earths ($\text{R}=\text{Tb}, \text{Dy}, \text{Ho}, \text{Yb}$) synthesized by rapid quenching of the liquid phase. The relations between the structures of the metastable and equilibrium phases and the transformations from the former to the latter are discussed by using X-ray and neutron diffraction, differential scanning calorimetry and magnetic measurements.

The magnetic and structural properties of RCu_5 with the cubic AuBe_5 -type structure have been investigated with the neutron diffraction and magnetic measurements [1]. In magnetisation measurements it has been found that TbCu_5 and DyCu_5 [2] behave antiferromagnetically below a temperature of 15 K and 7 K, respectively. For YbCu_5 sample the minimum at around 20 K in the temperature dependence of the Knight shift of ^{63}Cu at 4c site was shown [3], which may be associated with a formation of Kondo-lattice state.

The compound DyGe_6Mn_6 is known as a complicated helimagnet with hexagonal HfGe_6Fe_6 -type structure and it shows also an incommensurate magnetic order. Similar systems attracted much attention due to complicated magnetic ordering phenomena. Substitution of Fe instead Mn (ferromagnetic element instead antiferromagnetic element) in DyGe_6Mn_6 has been shown to influence the crystal and magnetic structures [4].

The point-contact-spectroscopy (PCS) enables us to gain energy spectra of quasiparticles like phonons, magnons or crystal-field excitations. Here, it was employed to study the impact of magnetic interactions on the electron-quasiparticle interactions in $\text{DyGe}_6\text{Mn}_{6-x}\text{Fe}_x$ for $x = 0, 3$ and 6. In the sample of pure DyGe_6Mn_6 we observe electron-phonon scattering and also scattering on magnetic quasiparticles. For all samples we observed a maximum in $dV/dI(V)$ around zero and an increase for voltages higher than 5 mV (for DyGe_6Fe_6) and 30 mV (for DyGe_6Mn_6). The point-contact characteristic of the compound with $x = 3$ shows a similar behaviour [5]. We suggest a magnetic origin of the maximum around zero voltage.

For comparison, the results of magnetic investigation obtained for DyFe_6Al_6 compound in the fully amorphous state will be discussed.

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