



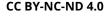
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## MAGNETIC ORDER IN THE HEAVY FERMION SYSTEM Ce(Cu<sub>1-x</sub>Ni<sub>x</sub>)<sub>2</sub>Ge<sub>2</sub>

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The magnetic phase diagram of the heavy fermion (HF) systems  $Ce(Cu_{1-x}Ni_x)_2Ge_2$  is discussed utilizing results of transport, thermodynamic and neutron-scattering measurements. While the Kondo temperature increases monotonically with x, a complex x-dependence is found for the Néel temperature, associated with a transition from local-moment to itinerant HF magnetism.

The tetragonal Ce-homologes of ThCr<sub>2</sub>Si<sub>2</sub>, while lacking direct 4f-wave function overlap, exhibit a distinct 4f-ligand hybridization. The latter, along with strong many-body renormalizations below the so called "lattice Kondo temperature" T\*, is effective in forming heavy-mass quasiparticles ("heavy fermions" (HF)). In the case of the compounds CeM2X2 (M: Cu, Ag, Au, Ru, Ni; X: Si, Ge), for large Ce-M distances r (≥ 3.3Å), magnetic ordering between local moments occurs at  $T_m$ , whereas in the limit of small r ( $\leq$  3.2Å) a valence fluctuating state forms [1]. In the intermediate regime, the HF-compounds CeNi<sub>2</sub>Ge<sub>2</sub>, CeRu<sub>2</sub>Si<sub>2</sub> and CeCu<sub>2</sub>Si<sub>2</sub> are found, of which the latter is unique not only because it is a superconductor below 0.7K [2], but also it seems to exhibit an itinerant type of HF magnetism below approximately 0.8K (at zero magnetic field) and 7 Tesla (T→0) [3].

In this communication, we report a brief summary of a systematic study on the quasibinary system  $Ce(Cu_{1-x}Ni_x)_2Ge_2$ , which reveals a transition from local-moment ordering to itinerant HF magnetism. These alloys are ideally suited for this purpose: The Kondo binding energy  $kT^*$  of  $CeCu_2Ge_2$  ( $T^*=8\pm 2K$ ) is of the same order of magnitude as the magnetic interaction energy,  $kT_{RKKY}$  ( $T_{RKKY}=7K$ ), at which short-range ordering sets in [4]. Long range magnetic order has been detected below  $T_m = T_{N1} = t^{-1} + t^{-1} +$ 

or magnitude as the magnetic interaction energy,  $KI_{RKKY}$  ( $T_{RKKY} = 7K$ ), at which short-range ordering sets in [4]. Long range magnetic order has been detected below  $T_m = T_{N1} = 4.1K$  [4,5]. On the other hand,  $CeNi_2Ge_2$  ( $T^* = 30K$ ) exhibits the signature of a Kondo lattice or HF compound with no long-range magnetic order [6]. For the samples prepared as described in [1], no parasitic phases have been detected by X-ray diffractometry or electron microprobe analysis.

Fig.1 shows the concentration dependence of the magnetic ordering temperature (in reduced units) for  $\text{Ce}(\text{Cu}_{1-x}\text{Ni}_x)_2\text{Ge}_2$  as determined from thermal expansion,

specific heat and magnetic susceptibility measurements [7,8]. Even small Ni concentrations cause a dramatic decrease of  $T_{N1}(x)$ , which extrapolates to zero near x=0.2. However, for larger x a second branch,  $T_{N2}(x)$ , develops with a maximum near x=0.5. This new magnetically ordered phase can be monitored up to approximately x=0.75. Bulk measurements show clear evidence for a second transition below  $T_{N1}$  and  $T_{N2}$ , respectively in the range  $0.02 \le x \le 0.3$  (hatched region) as discussed in [8].

The magnetic structure and the size of the ordered moments have been determined by neutron powder-diffraction experiments for the compounds with concentrations x=0, 0.1, 0.28 and 0.5. Incommensurate modulated spin arrangements with magnetic Bragg reflections  $|Q| = |\tau_{hkl} \pm q_0|$ , where (hkl) is a vector of the reciprocal chemical lattice and q<sub>n</sub> the propagation vector, have been found. They are described by the following values. For x=0:  $q_0 = (0.28, 0.28, 0.54)$  and the effective moment  $\mu_s = 0.74 \ \mu_B$ ; for x=0.1:  $q_0=(0.28, 0.28, 0.41), \mu_S=0.5\mu_B$ ; for x=0.28:  $q_0 = (0.11, 0.11, 0.25)$  and finally for  $x = 0.5 : q_0 = (0, 0, 0.14)$ ,  $\mu_{\rm S} = 0.3 \mu_{\rm R}$ . For this latter sample the magnetic Bragg reflections are much broader than the experimental resolution of the spectrometer which presumably reflects a disturbance of long-range magnetic order by single-site (Kondo) interactions.

To gain further insight into this competition of single-site and inter-site interactions, the magnetic relaxation rate via quasielastic neutron scattering has been investigated in the temperature range  $1.5K \le T \le 200K$ . The lattice Kondo temperature as a function of Ni concentration, as read off the residual quasielastic line widths at low temperatures ( $T \ge T_N$ ), is also plotted in Fig.1 and compared to  $T^*$  as determined from thermal expansion and resistivity

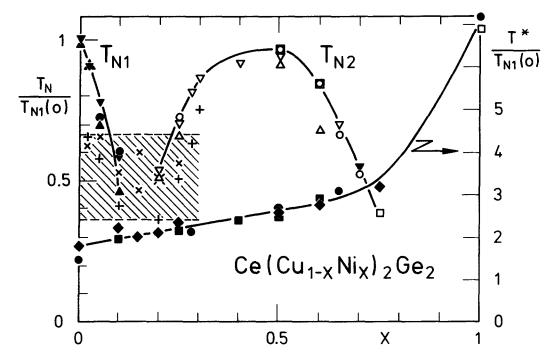


Fig.1. LEFT SCALE: x,  $T_N$  phase diagram for the quasi-binary compound  $Ce(Cu_{1-x}Ni_x)_2Ge_2$  as determined from specific heat  $(\P, \nabla, +)$ , thermal expansion  $(A, \Delta, x)$  and dc susceptibility  $(\P, O)$  measurements [9]. RIGHT SCALE:  $T^*$  as determined from the residual quasielastic neutron line width  $(\P)$ , thermal expansion  $(\P)$  and resistivity  $(\P)$  peaks. Positions of the latter are scaled by factors 1.5 and 1.9, respectively  $(T^*$  and  $T_N$  are normalized to  $T_{N,1}$  for x=0).

maxima. With increasing x,  $T^*$  increases continuously and, for x=0.65, the Kondo binding energy is found to be larger than the energy of the inter-site interactions by almost a factor of four.

The most exiting result of the present study is the occurence of an extremely short propagation vector for the x=0.5 sample which characterizes a static spin wave extending over almost ten lattice constants. Such short ordering wave vectors have been predicted by Grewe and Welslau [9], if HF band magnetism develops in a Kondo lattice. Recent calculations of Welslau and Grewe within this model [10] revealed the development of  $T_{\text{N}}$  and  $T^*$  as a function of the local exchange coupling constant  $g\!=\!N_{\text{F}}J$  ( $N_{\text{F}}$ : conduction band density of states at the Fermi level;  $J\!<\!0$ : exchange integral) in the full range between  $T_{\text{N}}\!\times\!T^*$  and  $T_{\text{N}}\!\times\!T^*$ . These calculations are corroborated by our results near  $x\!=\!0$  where  $T_{\text{N}}\!\approx\!T^*$  and for  $0.5\!<\!x\!\leq\!0.75$  where  $T_{\text{N}}\!\propto\!T^*$ .

The intermediate composition range, in which

- (i) two subsequent transitions are observed as a function of T for each sample  $(0.02 \le x \le 0.3)$  and
- (ii)  $T_{N2}(x)$  shows a monotonic increase (0.2  $x \le 0.5$ ),

challenges further investigation.

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