Tri-critical point and suppression of the Shastry-Sutherland phase in Ce$_2$Pd$_2$Sn by Ni doping

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Structural, magnetization and heat capacity measurements were performed on Ce$_2$(Pd$_{1-x}$Ni$_x$)$_2$Sn (0 ≤ x ≤ 0.25) alloys, covering the full range of the Mo$_2$FeB$_2$ structure stability. In this system, the two transitions observed in Ce$_2$Pd$_2$Sn (at $T_N = 4.8$ K and $T_C = 2.1$ K respectively) converge into a tri-critical point at $T_{cr} \approx 3.4$ K for $x \approx 0.3$, where the intermediate antiferromagnetic AF phase is suppressed. The $T_N(x)$ phase boundary decrease is due to an incipient Kondo screening of the Ce-4f moments and local atomic disorder in the alloy. Both mechanisms affect the formation of Ce-magnetic dimers on which the Shastry-Sutherland lattice (SSL) builds up. On the contrary, the $T_C(x)$ transition to the ferromagnetic ground state increases as a consequence of the weakening of the AF-SSL phase. Applied magnetic field also suppresses the AF phase like in the stoichiometric compound.

Keywords: Cerium compounds, Critical Points, Magnetic Phase Diagrams, Magnetic transitions

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I. INTRODUCTION

Crystalline structures with local symmetries which favor magnetic frustration attract special interest because they provide the scenario for novel phases formation. In many cases, magnetic frustration or exotic phases occur in competition with classical long range order ground states GS. Thus, the search of exotic phases is addressed to the vicinity of magnetic transitions, since the ‘roughness’ of the free energy may develop new relative minima as a function of a non trivial order parameter [1]. Those minima compete in energy for the formation of novel phases, which may become unstable under small variation of control parameters like magnetic field, alloying or pressure.

Beside the mentioned frustrated states, alternative phases may occur under peculiar geometrical conditions. Among them, the so called Shastry-Sutherland lattice (SSL) [2] [3] builds up from mutually orthogonal magnetic dimers which impose further topological and magnetic constrains. These conditions are realized in some members of the R$_2$T$_2$X family of compounds (with R = Rare Earths or Actinides, T = Transitions Metals and X = ‘p’ type metalloids), which crystallize in tetragonal Mo$_2$FeB$_2$ structure [4]. In that structure, each ‘R’ layer forms a mosaic of magnetic atoms coordinated as isoceles triangles between nearest- and next nearest neighbors centered on the z-axis of the T element. The shortest side of those triangles, where R-R magnetic dimers form, is shared by two consecutive triangles like in the shortest diagonal of a rhombohedron. The resulting simple square-lattice of mutually orthogonal rhombohedron mimicks a sort of ‘pinwheel’ centered on the z-axis of the X element [5], whereas the net of dimers form a simple two dimensional (2D) square lattice.

Recently, SSL phases were found in a number of R$_2$T$_2$X compounds [5] [6] and the involved magnetic interactions theoretically discussed [7]. In Ce$_2$Pd$_2$X compounds, where Ce-Ce dimers form due to a nearest neighbor ferromagnetic FM interaction, the SSL phase shows up within a limited range of temperature (c.f. between $T_N = 4.8$ K and $T_C = 2.1$ K in Ce$_2$Pd$_2$Sn [5]). Below $T_C$, a FM-GS takes over undergoing a first order transition due to the discontinuity in the order parameter. Further studies performed under magnetic field on the mentioned compound [6] showed that the intermediate phase is suppressed applying a magnetic field $B_{cr} \approx 0.12$ T at $T_{cr} \approx 3.2$ K. The fact that the SSL phase is suppressed by quite low magnetic field in Ce$_2$Pd$_2$Sn remarks its instability respect to a 3D FM magnetic structure as GS.

In this work we have investigated Ni doped Ce$_2$(Pd$_{1-x}$Ni$_x$)$_2$Sn alloys with the scope to compare the effect of structural pressure and magnetic field on the stability of the SSL phase. Since Pd and Ni are iso-electronic elements, but being Ni atoms about 25% smaller in volume than Pd ones, an effective structural pressure is expected to weak the Ce-4f magnetic moments. Similarly, applied magnetic field shall progressively suppress the SSL phase like in the stoichiometric compound.

In a former study performed on this system [9], no change of structure was reported under Ni doping. The unit cell volume was reported to follow a Vergard’s law up to $x = 0.6$, with the paramagnetic Curie-Weiss temperature $\theta_P$ decreasing from 20 K at $x = 0$ to -40 K at $x = 0.5$. Electrical resistivity ($\rho$) was observed to be nearly temperature independent with a high value of $\rho_0$ and large difference upon cooling and heating, which was interpreted as due to cracks in the alloyed samples [9]. However, in a recent investigation, a change of structure was detected at $x = 0.35 \pm 0.05$ [10]. Thus, the large $\rho_0$ values and the abnormal $\rho(T)$ dependence can be explained as due to non single crystalline phases in the
concentration range where the system presents a coexistence of two crystalline structures.

II. EXPERIMENTAL DETAILS AND RESULTS

Details for sample preparation were described in a previous paper [5]. Structural characterization confirms the single phase composition of the samples in a tetragonal Mo$_2$FeB$_2$-type structure for $x < 0.3$. Beyond a short range of coexistence of two phases, this system stabilizes in an orthorhombic W$_2$CoB$_2$ type structure for $x > 0.4$ [10]. The actual composition of the stoichiometric compound was determined to be Ce$_{2,005}$Pd$_{1,985}$Sn$_{0,997}$ after SEM/EDAX analysis. Lattice parameters slightly decrease with Ni concentration from $a = 7.765\text{Å}$ and $c = 3.902\text{Å}$ at $x = 0$, to $a = 7.7122\text{Å}$ and $c = 3.8941\text{Å}$ at $x = 0.25$. These variations drive a reduction of the unit cell volume of about $\approx 1.5\%$ between $x = 0$ and 0.25. On the contrary, the ‘c/a’ ratio practically does not change, indicating that the local symmetry is not affected by doping.

DC-magnetization measurements were carried out using a standard SQUID magnetometer operating between 2 and 300 K, and as a function of field up to 5T. Specific heat was measured using standard heat pulse technique in a semi-adiabatic He-3 calorimeter in the range between 0.5 and 20 K, at zero and applied magnetic field up to 4T. The magnetic contribution to the specific heat $C_m$ was computed subtracting the phonon contribution extracted from the isotypic La$_2$Pd$_2$Sn compound. Electrical resistivity was measured between 0.5 K and room temperature using a standard four probe technique with an LR700 bridge. However, sample cracking inhibits to extract valuable information from $\rho(T)$ data.

As shown in Fig. 1, Ni doping does not affect significantly the magnetic susceptibility ($\chi$) at high temperature. Fits of $\chi(T)$ do not show significant variation of crystal field effect (CEF) from the values observed in stoichiometric Ce$_2$Pd$_2$Sn [5], with respective splitting at $\Delta_I = 65 \pm 5$ K and $\Delta_{II} = 240 \pm 10$ K. From the inverse of $\chi(T)$, one extracts that the high temperature $\theta_{HF}^{HT}$ slightly increases only for $x \geq 0.2$ from -16 K to $\approx 25$ K. This indicates that only a minor hybridization increase occurs for the CEF excited levels within this range of Ni doping. Coincidentally, the high temperature Ce magnetic moment practically does not change from its Ce$^{3+}$ value.

A different behavior is observed at low temperature, where $\theta_{LT}^{HF} > 0$. Its value decreases from 4.5 K (at $x = 0$) for $x = 0.1$.

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**FIG. 1.** (Color online) High temperature magnetic susceptibility in a Log-Log representation for different Ni concentrations (left and upper axes) and inverse susceptibility (right and lower axes).

**FIG. 2.** (Color online) (a) Low temperature inverse magnetic susceptibility showing the variation of $\theta_{HF}^{HT}$, measured with $B = 0.1$ T. (b) Field magnetization within the FM phase, showing the decrease of the saturation moment with Ni concentration increase. Inset: magnetic moment at $B = 5$ T within the $x \leq 0.25$ range.
to 3.7 K (at $x = 0.25$), as shown in Fig. 2b, indicating a moderate Ni doping effect on the doublet GS magnetism. This goes hand by hand with the magnetic saturation extracted from field dependent magnetization measurements (see Fig. 2b), performed at $T = 1.8$ K within the FM phase. Also the cusp of $M(T)$ at $T = T_N$ is reduced till it is overcome by the FM contribution (see Fig. 2b).

In the case of specific heat, Ni concentration affects quite significantly the $C_m(T)$ jumps of both transitions as shown in Fig. 3. While the temperature of the upper one decreases from $T_N(x = 0) = 4.8$ K to $T_N(x = 0.25) \approx 3.5$ K, the lower transition increases from $T_C(x = 0) = 2.1$ K up to $T_C(x = 0.25) \approx 3.3$ K (see inset in Fig. 3). These opposite Ni dependencies drive the transitions to merge at a tri-critical point at $T_{cr} \approx 3.4$ K, just above the highest Ni concentration studied ($x = 0.25$).

$T_N(x)$ is a second order transition, whereas $T_C(x)$ is of first order. This character is recognized by an hysteretic $C_m(T)$ dependence measured around $T_C$ and reflects the discontinuity in the magnetic order parameter at $T = T_C$. Both $C_m(T)$ maxima decrease with Ni doping, however as they merge into an unique maximum the $C_m(T)$ cusp clearly rises. Notice the sharpness of the cusp in sample $x = 0.25$ revealing its vicinity to the critical point.

### B. Magnetic field effects

Doping and magnetic field effects coincide enhancing the FM character of the GS phase with the consequent $T_C(x, B)$ increase. On the contrary, while Ni doping produces a decrease of $T_N(x)$ magnetic field practically does not affect the upper transition. The doping effect can be observed in Fig. 4a, which shows how the temperature of the spontaneous magnetization (related to $T_C$) increases, overlapping the $M(T_N)$ cusp as Ni content increases.

In the case of applied magnetic field, one can observe in Fig. 4b for sample $x = 0.1$ how the temperature of the $M(T_N)$ cusp remains practically unchanged. The same behavior was observed in the stoichiometric compound $[8]$ and is confirmed in sample $x = 0.2$. In the case of sample $x = 0.25$, the $T_N$ cusp is already overlapped by the FM signal at very low applied field. One have to remind that, apart from the $T_C(x)$ increase, the FM transition is smeared under applied magnetic field and therefore the high temperature tail of the FM magnetization overcomes the $T_N$ cusp faster than the actual increase of $T_C(x)$.
III. DISCUSSION

A. Specific heat Gap

Relevant information related to the magnetic structure of this system can be extracted from the analysis of the thermal dependence of the specific heat within the ordered phase. Since the $x = 0$ sample has shown strong anisotropic effects, we have analyzed the magnetic contribution of the FM phase to $C_m(T)$ by fitting all curves with an unique function in order to evaluate the variation of the characteristic parameters as a function of Ni content. The simplest applicable function is $C_m / T = \gamma + AT^2 \exp(-E_g / T)$, where the $\gamma$ term accounts for the degrees of freedom behaving as a Kondo liquid and $E_g$ represents the energy (expressed in temperature) of the gap of anisotropy in the magnon spectrum. Respective fits are included in Fig. 5 using a double logarithmic scale, and computed values are collected in the inset. These results indicate that the specific heat behaves as that of systems whose gap of anisotropy deceases to zero when approaching the critical concentration. Coincidentally, the $\gamma$ contribution increases with Ni content.

B. Entropy and Kondo Temperature

From the thermal variation of the magnetic contribution to the entropy ($S_m$), the effect of Ni doping on the Ce-4f magnetic moment can be traced. As it can be observed in Fig. 6 it is only for $x \geq 0.2$ that some incipient hybridization effect can be detected. To extract the Kondo temperature variation, we have applied the current Desgranges-Schotte criterion \[11\] from which at $T = T_K$ the entropy reaches the value $S_m = 0.66 \ln 2$ value from which $T_K$ is extracted.

C. Stability of the SSL phase versus doping

In order to investigate up to which extend Ni doping affects the stability of the SSL phase, we have studied the magnetic field effect in the alloyed samples follow-
FIG. 7. (Color online) (a) Field dependence of magnetization of sample $x = 0.1$ between 3 and 4.2 K. (b) First derivative and (c) second derivative for the $3.7 \leq T \leq 4.1$ K range, with the arrow indicating the satellite anomaly.

FIG. 8. (Color online) Ni concentration and magnetic field dependent phase diagram in a 3D representation.

The comparison between doping and magnetic field effects is presented in a 3D phase diagram in Fig. 8. There, the field driven transition between intermediate AF and FM-GS phases is drawn according to the temperature of the maximum slope of $M(T)$ (i.e. $\partial M/\partial T |_{_{max}}$) which is in agreement with $C_{m}(T)$ results. Both phase boundaries join at a critical point, which decreases with Ni content from $T_{cr}(x = 0) = (4.2 \pm 0.3)$ K and $B_{cr} = (0.16 \pm 0.02)$ T to $T_{cr}(x = 0.25) \approx (3.5 \pm 0.3)$ K and $B_{cr} \approx (0.02 \pm 0.005)$ T, at the edge of the intermediate phase disappearance (i.e. where $B_{cr} \rightarrow 0$). Unfortunately, the tri-critical point where AF, FM and paramagnetic phases join together cannot be reached by doping because it lies beyond the limit of stability of the respective curves are collected in Fig. 7c, providing better information to determine the onset of the induced FM phase (main maximum at $B \approx 0.077$ T) and the satellite anomaly (see the arrow at $B \approx 0.12$ T).

In comparison with the stoichiometric compound, the main maximum has decreased from $B \approx 0.11$ to 0.08 T, whereas the satellite shifted from $B \approx 0.15$ to 0.12 T with a significant reduction in the intensity. Although the $B$ values follow the predictions for a SSL for the appearance of a plateaux in the magnetization (at 1/4 and 1/8 of the saturated moment $[3]$), these results from sample $x = 0.1$ are in the limit of our experimental detection. Effectively, in the following concentration studied ($x = 0.2$) no traces of the mentioned anomaly were observed indicating a border line for the SSL formation at $x \approx 0.1$. Not only the weakening of Ce magnetic moments but also the local disorder due to the difference between Pd and Ni sizes inhibit the Ce-Ce dimers formation with the consequent smearing of the SSL phase.

D. Magnetic phase diagram

The comparison between doping and magnetic field effects is presented in a 3D phase diagram in Fig. 8. There, the field driven transition between intermediate AF and FM-GS phases is drawn according to the temperature of the maximum slope of $M(T)$ (i.e. $\partial M/\partial T |_{_{max}}$) which is in agreement with $C_{m}(T)$ results. Both phase boundaries join at a critical point, which decreases with Ni content from $T_{cr}(x = 0) = (4.2 \pm 0.3)$ K and $B_{cr} = (0.16 \pm 0.02)$ T to $T_{cr}(x = 0.25) \approx (3.5 \pm 0.3)$ K and $B_{cr} \approx (0.02 \pm 0.005)$ T, at the edge of the intermediate phase disappearance (i.e. where $B_{cr} \rightarrow 0$). Unfortunately, the tri-critical point where AF, FM and paramagnetic phases join together cannot be reached by doping because it lies beyond the limit of stability of the respective curves are collected in Fig. 7c, providing better information to determine the onset of the induced FM phase (main maximum at $B \approx 0.077$ T) and the satellite anomaly (see the arrow at $B \approx 0.12$ T).
The evolution of AF and FM phase boundaries respond to different effects. Starting from high temperature, the decrease of the upper (AF) transition $T_N(x)$ is expected from i) the competition between Kondo effect and RKKY magnetic interactions according to Doniach-Lavagna model [12] acting on the Ce-4f magnetic moments intensity and ii) the local atomic disorder introduced by alloying atoms of different sizes. Both effects weaken the formation of magnetic Ce-dimers on which the SSL builds up. On the contrary the $T_C(x)$ increase is a consequence of the intermediate AF phase weakening like in Ce$_2$Pd$_2$Sn under applied magnetic field [8].

The effect of magnetic field is qualitatively similar for each concentration accounting that the range of stability of the AF phase is progressively reduced. Notably, the $\partial M/\partial B \mid_T$ derivatives of sample $x = 0.25$ still display a weak maximum in a restricted range of field and temperature. This indicates that the critical Ni concentration is not reached yet despite in the $C_m(T)$ cusp of that sample both phase boundaries seem to have converged. Interestingly, the critical temperature $T_{Cr}$ reached by doping at zero field and by magnetic field applied on the stoichiometric compound, and allows to better define the tri-critical point extrapolated at $T_{Cr} \approx 3.4$ K and $x = 0.3$, where $B_{cr} \to 0$. Unfortunately $x = 0.3$ lies within the structural instability range. Nevertheless, further attempts to get closer to that critical point, and to investigate the Ni-rich side of the phase diagram are in progress.

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