Magnetic behaviour of the new hydride CePtAlH_x

Jean-Louis Bobet, Bernard Chevalier, François Weill, Jean Etourneau

► To cite this version:

Jean-Louis Bobet, Bernard Chevalier, François Weill, Jean Etourneau. Magnetic behaviour of the new hydride CePtAlH_x. Journal of Alloys and Compounds, 2002, 330-332, pp.373-375. 10.1016/S0925-8388(01)01578-X . hal-00070926

HAL Id: hal-00070926

<https://hal.science/hal-00070926>

Submitted on 30 May 2012

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Magnetic behaviour of the new hydride CePtAlH_x

Bobet J.-L., Chevalier B., Weill F., Etourneau J.

Published in Journal of Alloys and Compounds 2002, vol. 330-332, p. 373-375.

Abstract :

CePtAl absorbs hydrogen up to a concentration of $1.1(1)\text{H mol}^{-1}$ at room temperature. The formation of this hydride CePtAlH_{1.1} that loses hydrogen in air corresponds to a plateau pressure of 0.08–0.1 MPa. Its absorption–desorption is practically reversible which can present a potential interest for applications (but only a few due to the high price of the material) around room temperature and normal pressure. Hydrogenation of CePtAl induces both: (i) a structural transition but the structure of the hydride is unknown at present; (ii) an increase of its ferromagnetic character; the hydride has a Curie temperature $T_c=11.6(2)$ K twice greater than that determined for the initial intermetallic (i.e. $5.6(2)$ K).

1. Introduction

Recently, we have shown that the hydrogenation of the CeNiAl compound induces both a structural transition from the hexagonal ZrNiAl-type to the hexagonal AlB₂-type for the hydride CeNiAlH_{1.93} and a valence transition for cerium from tetravalent to trivalent state [1]. The hydrogenation of CeNiAl leads to an expansion of its molar volume favouring the occurrence of spin fluctuation behaviour below 7 K for the hydride. Encouraged by this interesting result, we are now studying the hydrogen absorption–desorption properties of the equiatomic CePtAl compound. This ternary aluminide crystallises in the orthorhombic TiNiSi-type structure where Ce-atoms form a network of chains parallel to the *a*-axis [2]. Neutron powder diffraction experiments performed on CePtAl reveal a complex magnetic phase diagram: three successive transitions are detected respectively at $T_c=5.9$ K, $T_2=4.3$ K and $T_3=2.2$ K [3]. But this study shows that the magnetism of CePtAl is dominated by a ferromagnetic component $1.35(4) \mu_B$ at 1.5 K of the ordered Ce-moment. This original behaviour results from the existence of a large magnetic anisotropy caused by competition between magnetic exchange interactions existing respectively in the Ce–Ce chains and between themselves [4]. Moreover, specific heat measurements suggest that the reduction of the ordered Ce-moment found in CePtAl can be explained by the existence of the Kondo effect [5]. The magnetic and Kondo interactions are governed by the Ce–Ce interatomic distances ($d_{\text{Ce-Ce}}$); in this view it is interesting to modify these distances for instance by insertion of hydrogen in the lattice.

In this paper, we report on the synthesis of the hydride CePtAlH_x and on its characterisation using X-ray powder diffraction and magnetization measurement.

2. Experimental details

CePtAl sample was prepared by arc-melting stoichiometric amounts of pure elements (3N for Ce, 3N5 for Pt and 5N for Al) under a protective Ar-atmosphere followed by an annealing treatment under vacuum at 1073 K for 1 month. Its characterisation by electron microprobe analysis reveals a perfect chemical homogeneity. Its checking by X-ray powder diffraction (Philips PW 1050 diffractometer, Cu-K α radiation) confirms that CePtAl crystallises in the orthorhombic TiNiSi-type with unit cell parameters $a=0.7208(5)$ nm, $b=0.4481(5)$ nm and $c=0.7740(5)$ nm in agreement with those reported previously [2].

Hydrogen absorption–desorption measurements were performed using the apparatus described previously [6]. CePtAl sample was crushed (grain size $\leq 100 \mu\text{m}$), and then exposed to hydrogen gas. The amount of H-atom absorbed was determined volumetrically by monitoring pressure changes in a calibrated volume.

Magnetization measurements were performed using a Superconducting Quantum Interference Device (SQUID) magnetometer in the temperature range 1.8–300 K and applied fields up to 5 T.

3. Results and discussion

Without any thermal activation treatment, CePtAl absorbs hydrogen at room temperature. Its hydrogen absorption–desorption capacities are strongly dependent to the number of hydriding–dehydriding cycles (Fig. 1): (i) at the initial cycle $n=1$, CePtAl starts to absorb hydrogen at a pressure of 0.8 MPa; (ii) for the second one, the amount of H-atom absorbed varies almost linearly with the hydrogen pressure; (iii) finally, one can observe that the sample is successfully activated after three hydriding–dehydriding cycles. At 295 K, CePtAl absorbs hydrogen up to a concentration of $1.1(1) \text{ H mol}^{-1}$ under a pressure of 1.1 MPa (Fig. 2). The plateau pressure at this temperature in the range 0.08–0.1 MPa corresponds to the formation of the hydride $\text{CePtAlH}_{1.1(1)}$ that loses hydrogen in air (as reported on Fig. 3). Its absorption–desorption is practically reversible.

Pressure–composition–temperature (PCT) curves realized at various temperatures for CePtAl are presented on Fig. 2. For this study, the sample was hydrided–dehydrided three times before the shown experiments were carried out; so these results are expected to represent the thermodynamically stable state. It appears clearly that the increase of temperature induces a significant decrease of the hydrogen content of the CePtAlH_x hydride ($x \approx 0.9(1)$ at 373 K) and increase of the plateau pressure. The rather large plateau slope observed at 373 K can be attributed to sample temperature inhomogeneities. The thermodynamic data calculated according to the Van't Hoff equation are $\Delta H = -24 \text{ kJ mol}^{-1}$ and $\Delta S = 180 \text{ J mol}^{-1} \text{ K}^{-1}$.

The X-ray powder pattern of CePtAl, given in Fig. 3, is completely indexed on the basis of the orthorhombic TiNiSi -type structure. On the contrary the pattern relative to $\text{CePtAlH}_{1.1}$ hydride, characterising a well-crystallised material, clearly shows that hydrogenation leads to a structural transition (Fig. 3). This last structure is unknown at present but under study using transmission electron microscopy. After exposure in air for 5 days at room temperature, the hydride $\text{CePtAlH}_{1.1}$ is partially decomposed; X-ray diffraction analysis reveals the occurrence of a mixture containing CePtAl and $\text{CePtAlH}_{1.1}$ (Fig. 3). For instance, the diffraction peaks (111) and (103) relative to CePtAl compound are clearly observed after the exposure. This result agrees with the low dissociation pressure $\approx 0.1 \text{ MPa}$ at 295 K (Fig. 2) of this hydride. After desorption (0.05 MPa for 1 h) the X-ray diffraction analysis reveals the occurrence of a single phase which is similar to CePtAl (as expected).

Magnetization measurements were performed on the hydride $\text{CePtAlH}_{1.1}$ powder enclosed in epoxy resin in order to prevent the hydrogen loss. Above 70 K, the reciprocal magnetic susceptibility χ_m^{-1} of this hydride follows a Curie–Weiss law $\chi_m^{-1} = (T - \theta_p)/C_m$ having as Curie constant $C_m = 0.65 \text{ emu mol}^{-1}$ and a large and negative paramagnetic Curie temperature $\theta_p = -44 \text{ K}$. The effective magnetic moment $\mu_{\text{eff.}} = (8C_m)^{1/2} = 2.3 \mu_B/\text{Ce-atom}$ is smaller than that calculated for free Ce^{3+} ion ($2.54 \mu_B$). Similar investigation performed on CePtAl (before any treatment or even after hydrogen desorption) and gives $\mu_{\text{eff.}} = 2.51 \mu_B/\text{Ce-atom}$ and $\theta_p = -37 \text{ K}$.

The temperature dependence below 17 K of the magnetization of CePtAl and $\text{CePtAlH}_{1.1}$ is shown in Fig. 4. The curve obtained for CePtAl is similar to that observed previously by Kitazawa et al. [5]; the curve exhibits a strong increase at $T_c = 5.8(2) \text{ K}$. A similar behaviour is observed at $T_c = 11.6(2) \text{ K}$ for $\text{CePtAlH}_{1.1}$ hydride. In other words, the hydride has a Curie temperature twice greater than that determined for CePtAl. This fact is confirmed by the curves showing at low temperatures, the field dependence of the magnetization (Fig. 5); these are typical for a ferro- or ferrimagnetic material. At 4 K, the remanence is $0.15 \mu_B \text{ mol}^{-1}$ and the magnetization takes a low value of $0.5 \mu_B \text{ mol}^{-1}$ in applied field $B = 3.5 \text{ T}$. This last value is weak compared to that expected $g_J J = 2.14 \mu_B \text{ mol}^{-1}$ for Ce^{3+} ion which suggests that the magnetic ordering of $\text{CePtAlH}_{1.1}$ is not purely ferromagnetic.

4. Conclusion

In conclusion, our study shows that CePtAl absorbs and desorbs hydrogen reversibly at 295 K. This presents a potential interest for application around ambient temperature and pressure. The insertion of hydrogen induces both a structural transition and a strong increase of its ferro (ferri)magnetic character.

References

1. J.-L. Bobet, B. Chevalier, B. Darriet, M. Nakhl, F. Weill, J. Etourneau, J. Alloys Comp., 317–318 (2001), p. 67.
2. F. Hulliger, J. Alloys Comp., 196 (1993), p. 225.
3. A. Dönni, H. Kitazawa, P. Fischer, J. Tang, M. Kohgi, Y. Endoh, Y. Morii, J. Phys: Condens. Mat., 7 (1995), p. 1663.
4. H. Kitazawa, S. Nimori, J. Tang, F. Iga, A. Dönni, T. Matsumoto, G. Kido, Physica B, 237–238 (1997), p. 212.
5. H. Kitazawa, A. Dönni, L. Keller, J. Tang, F. Fauth, G. Kido, J. Solid State Chem., 140 (1998), p. 233.
6. J.-L. Bobet, S. Pechev, B. Chevalier, B. Darriet, J. Alloys Comp., 267 (1998), p. 136.

Fig. 1. Pression-composition curves for CePtAl at 295 K and various hydriding-dehydriding cycles (n).

Fig. 2. Pression-composition curves for CePtAl at 295, 323 and 373 K.

Fig. 3. X-ray powder pattern of CePtAl and CePtAlH_{1.1} before and after exposure in air for 5 days.

Fig. 4. Temperature dependence of magnetization, measured in an applied field of $B=0.05$ T, for CePtAl and CePtAlH_{1.1}.

Fig. 5. Field dependence of magnetization for $\text{CePtAlH}_{1.1}$ measured at various temperatures.