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Crystal and magnetic structures of $LaNi_{5-x}Mn_x$

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The crystallographic and magnetic properties of $\text{LaNi}_{5-x}\text{Mn}_x$ with x=1, 1.5, and 2, have been investigated using x-ray and neutron diffraction, vibrating sample magnetometer and superconducting quantum interference device measurements. All the samples crystallize in the hexagonal CaCu₅-type (*P*6/mmm) structure. Manganese atoms preferentially occupy the 3*g* site in the LaNi₅ structure. The bulk magnetization of the LaNi_{5-x}Mn_x powder samples decreases rapidly as nickel is replaced by manganese. A ferrimagnetic model is applied to describe the magnetic structure of the LaNi_{5-x}Mn_x samples for x=1.5 and 2. A ferromagnetic model gives the best fit of the neutron diffraction data for the LaNi₄Mn sample. © 2000 American Institute of Physics. [S0021-8979(00)77408-1]

I. INTRODUCTION

Even though rare earth-transition-metal (R-M) intermetallics are the leading candidates for the next-generation permanent magnets, the relationship between the electronic structure and the magnetic properties of these compounds has not been fully understood. An explanation based on electronic structure as to why negative magnetic interactions are more common in R-Mn intermetallics than in R-Fe intermetallics¹ will be helpful in engineering the next generation of commercial permanent magnets. We intend to make use of recent developments² in spin-resolved photoemission spectroscopy to study in detail the band structure of these intermetallics. Because the interpretation of spinresolved photoemission spectra of complex intermetallics such as RM₁₂ and R₂M₁₇ is very difficult, it is prudent to start with the relatively simple RM5 system. However, because RFe5 and RMn5 intermetallics do not exist, we will study the electronic structures of the $LaNi_{5-x}Fe_x$ and LaNi_{5-x}Mn_x systems.

A good understanding of the crystallographic and magnetic properties of these intermetallics is necessary to accurately interpret their spin-resolved photoemission spectra. The LaNi_{5-x}Fe_x system has been investigated in detail in previous studies.^{3–5} These studies suggested the coexistence of long-range ferromagnetic ordering and spin-glass-like localized clusters. The crystallographic properties of LaNi_{5-x}Mn_x at 295 K have also been studied^{6,7} using x-ray and neutron diffraction. In the present study, the magnetic structures and properties of LaNi_{5-x}Mn_x are investigated using low-temperature neutron diffraction, superconducting quantum interference device (SQUID), and vibrating sample magnetometer (VSM) measurements, and are compared with those properties of LaNi_{5-x}Fe_x.³

II. EXPERIMENT

The LaNi_{5-x}Mn_x samples were prepared from elements of 99.99% purity or higher by induction melting in a cold copper crucible. Induction-melted ingots were annealed for two weeks at 750 °C. The phase purity of the samples was checked by x-ray diffraction with $Cu K \alpha$ radiation on a Scintag XDS 2000 x-ray diffractometer equipped with a singlecrystal graphite monochromator. The powder neutron diffraction data were obtained at the University of Missouri Research Reactor using neutrons of wavelength 1.4875 Å. The samples were placed in thin-walled vanadium containers, and neutron diffraction data were collected at 20 and 295 K for 4-6 h. The diffraction data were refined by the Rietveld technique.8 The bulk magnetic properties of the powder samples were measured at Southern Illinois University-Carbondale on a Quantum Design SQUID magnetometer, and at the Laboratoire des Materiaux et du Genie Physique (LMGP) in France on an Oxford VSM or Foner-type magnetometer.

III. RESULTS AND DISCUSSIONS

The LaNi_{5-x}Mn_x samples crystallized in the CaCu₅-type structure (space- group *P6*/mmm) with little (<5 at. %) or no impurities. Table I gives the lattice parameters and site occupancies of LaNi_{5-x}Mn_x compounds obtained from Rietveld refinements of the neutron diffraction data, and Fig. 1 shows the neutron diffraction data and fit for LaNi_{3.5}Mn_{1.5} at 20 K. The weak peak at $2\theta \approx 23^{\circ}$ is due to a nonmagnetic impurity. The unit cell expands with increasing manganese content at a rate of ~5.5% per substituted atom. This expansion rate is more than twice the rate of 2.5% per substituted atom observed in LaNi_{5-x}Fe_x, ³ which also crystallized in the CaCu₅-type structure. The difference in expansion rates

TABLE I. Lattice parameters, unit-cell volumes, and manganese site occupancies for $LaNi_{5-x}Mn_x$.

	295 K			20 K			% Mn occ.	
	a (Å)	<i>c</i> (Å)	$V(Å^3)$	a (Å)	<i>c</i> (Å)	$V(\text{\AA}^3)$	3 <i>g</i>	2 <i>c</i>
LaNi ₅	5.0162(1)	3.9837(1)	86.812(3)					
LaNi ₄ Mn	5.0940(2)	4.0697(2)	91.456(7)	5.0690(2)	4.0527(2)	90.182(7)	29.4(3)	5.2(5)
LaNi3.5Mn1.5	5.1392(2)	4.1104(2)	94.017(7)	5.1147(1)	4.0966(1)	92.810(3)	45.1(3)	6.1(6)
LaNi ₃ Mn ₂	5.1941(2)	4.1408(2)	96.746(7)	5.1741(2)	4.1329(2)	95.819(7)	60.8(3)	7(1)

is most likely due to the size difference between iron and manganese atoms ($r_{\rm Fe}$ =1.27 Å and $r_{\rm Mn}$ =1.31 Å). The substitutional occupancy patterns in the two series of samples, however, are similar. As was the case for iron in LaNi_{5-x}Fe_x, manganese atoms in the LaNi_{5-x}Mn_x samples preferentially substitute for nickel atoms at the 3*g* site.

A comparison of the neutron diffraction patterns measured at 20 and 295 K, see Fig. 2, illustrates the effects of the magnetic structure on neutron scattering in $LaNi_{5-x}Mn_x$. The large intensities of the (100) and (101) reflections in the diffraction patterns measured at 20 K are due to magnetic

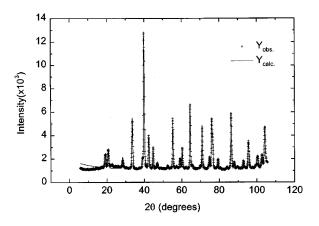


FIG. 1. Neutron diffraction pattern measured at 20 K for $LaNi_{3,5}Mn_{1,5}$ and a fit based on the CaCu₅-type structure and the ferrimagnetic mode.

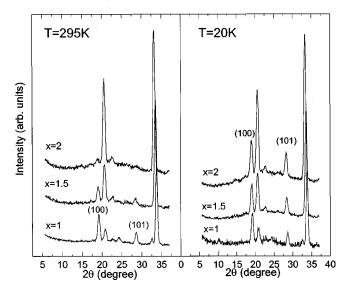


FIG. 2. Neutron diffraction patterns at 295 and 20 K for the $LaNi_{5-x}Mn_x$ samples. All peaks are normalized to the 111 reflections, which are not shown in the patterns.

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scattering. Furthermore, the difference between the intensities of these peaks at 20 and 295 K suggests that the atoms responsible for the observed magnetic scattering, predominantly the manganese atoms, possess appreciable magnetic moments.

Figure 3, which compares the magnetization versus temperature (*M* vs *T*) data for $LaNi_{5-x}Fe_x$ and $LaNi_{5-x}Mn_x$, illustrates the different effects of partially substituting the nickel sublattice with iron or manganese in LaNi₅. As seen in Fig. 3(a), partial substitution for nickel by iron increases the net magnetization of LaNi₅. Furthermore, the magnetic behavior of LaNi_{5-x}Fe_x samples with $x \ge 1.2$ is indicative of ferromagnetic ordering.⁹ In contrast, the net magnetization of LaNi_{5-r}Mn_r intermetallics decreases with increasing manganese content. The shapes of the M vs T curves and the weak magnetization values (<0.7 emu/g) for LaNi_{5-r}Mn_r intermetallics are not those of typical ferromagnetically ordered materials. In addition, these intermetallics do not reach magnetic saturation even at applied fields as high as 15 T.¹⁰ These observations, when combined with the fact that manganese atoms possess non-negligible magnetic moments, point to a magnetic structure consisting of a ferrimagnetically ordered manganese sublattice. As described below, detailed analysis of neutron diffraction data measured at 20 K confirms the existence of such a magnetic structure in the LaNi_{5-x}Mn_x intermetallics.

To simplify the analysis of the magnetic scattering contribution to the neutron diffraction data, the magnetic moments on the 2c manganese atoms were neglected due to the very small manganese occupancy of that site. Based on this assumption, the manganese atoms at the 3g sites are mainly responsible for the magnetic effects. Because all the magnetic reflections are allowed reflections of the P6/mmm space group, the crystallographic and magnetic unit cells are the same. Due to the observed weak magnetization values and the strong magnetic reflections, refinements of the 20 K neutron diffraction data of the LaNi_{5-x}Mn_x samples with x= 1.5 and 2.0 were done with a ferrimagnetic model. In this model, the three crystallographically identical 3g sites occupied by the manganese atoms, labeled 3g-1, 3g-2, and 3g-3in Fig. 4, are magnetically nonequivalent. The best fits of the neutron diffraction pattern were obtained when it is assumed that the magnetic moment of the manganese atom at the 3g-1 site is antiparallel to those at the 3g-2 and 3g-3 sites. This magnetic structure can be considered as a stacking of ferromagnetic sheets, where the magnetic moments are aligned parallel within each vertical sheet but alternate in direction in adjacent layers. In order to obtain better fits,

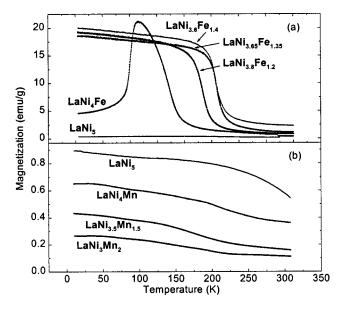


FIG. 3. Magnetization vs temperature for (a) $\text{LaNi}_{3-x}\text{Fe}_x(H=1000 \text{ Oe})$ and (b) $\text{LaNi}_{5-x}\text{Mn}_x(H=2000 \text{ Oe})$.

weak moments (0.1 μ_B or less) on the 2*c* and 3*g* nickel atoms were included in the refinements. The nickel moments, which might be induced by the manganese moments, are parallel to the net magnetic moment of the manganese sublattice. These weak moments could also be an artifact resulting from the neglected manganese moments at the 2*c* site, although they are similar to the induced moments found in LaNi_{5-x}Fe_x.^{3,4} The fit shown in Fig. 1 was obtained using this ferrimagnetic model.

For the LaNi₄Mn sample, however, a ferromagnetic model in which the magnetic moments of all manganese atoms are parallel gives the best fit of the neutron diffraction pattern at 20 K. Probability calculations show that the

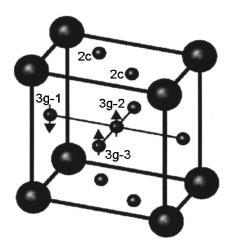


FIG. 4. LaNi_{5-x}Mn_x structure showing the antiparallel coupling of the manganese 3g moments. The larger atoms are La and the smaller ones are either Ni or Mn.

TABLE II. Total magnetic moments for $LaNi_{5-x}Mn_x$.

	$\mu_{ ext{total}}$ ($\mu_B/ ext{f.u.}$)	θ^{a} (degrees)
LaNi ₄ Mn	1.59	0
LaNi _{3.5} Mn _{1.5}	1.11	90
LaNi ₃ Mn ₂	1.07	115

^aAngle between the z axis and moment direction.

LaNi₄Mn compound has only 0.68 Mn-Mn nearest-neighbor pairs at the 3g site per unit cell. It means that the chance of having negative exchange (Mn–Mn antiparallel coupling) at the 3g site is less than 23% (0.68: $3 \approx 0.226$: 1, where 3 is the maximum possible number of nearest-neighbor pairs at the 3g site), which explains the ferromagnetic behavior of the LaNi₄Mn sample due to the dilute manganese concentration. The magnetic moments of the 3g manganese atoms obtained from the refinements are 1.25, 1.50, and 1.20 μ_B for the x =1, 1.5, and 2 samples. Table II gives the magnitude and direction of the total magnetic moments calculated from the neutron data refinements. The refined results suggest the rotation of the total moment direction away from the z axis to the xz plane when the manganese content increases. The total moment decreases with manganese content in agreement with the M vs T data [Fig. 3(b)].

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