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Mahmud Khan Southern Illinois University Carbondale

Shane Stadler Southern Illinois University Carbondale

Naushad Ali Southern Illinois University Carbondale

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Magnetocaloric properties of Fe and Ge doped Ni₂Mn_{1-x}Cu_xGa

Mahmud Khan,^{a)} Shane Stadler, and Naushad Ali

Department of Physics, Southern Illinois University, Carbondale, Illinois 62901

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The magnetocaloric properties of Fe and Ge doped Ni₂Mn_{0.75}Cu_{0.25}Ga Heusler alloys have been investigated. Using Ni₂Mn_{0.75}Cu_{0.25}Ga as the parent material, the Fe doped system $(Ni_2Mn_{1-x}(Cu-Fe)_xGa)$ and a Ge doped system $(Ni_2Mn_{1-x}Cu_xGa_{1-x}Ge_x)$ were studied. The manipulation of the Mn–Cu subsystem with Fe doping results in a decrease of the first order magnetostructural transition temperature, whereas the substitution of Ge for the Mn–Cu–Ga subsystems results in an increase of the magnetostructural transition temperature. In both cases the giant magnetocaloric effect is successfully preserved. © 2007 American Institute of Physics. [DOI: 10.1063/1.2712304]

Recently many ferromagnetic materials have been discovered that undergo first order magnetic transitions and exhibit large magnetocaloric effects (MCE).¹⁻³ Due to these recent discoveries, interest in the magnetocaloric cooling technology has grown significantly. When compared to the currently employed gas cooling technology, the magnetocaloric cooling technology has considerably enhanced efficiency, and therefore the recent discoveries and continuing research on magnetocaloric materials may lead to intense universal consequences.^{4–7} The MCE is a result of the alignment of magnetic moments with an external magnetic field. The alignment causes a reduction in the magnetic randomness or the magnetic component of the total entropy. The reduction of magnetic entropy is compensated by an increase in the other components of the total entropy. In the case of magnetocaloric materials, the compensation results in the heating of the material. A detailed discussion of the thermodynamics of the MCE is presented in Ref. 8.

The Heusler alloy Ni₂MnGa is a well known shape memory alloy that has potential application as a magnetic actuator material. Lately it has gained additional interest due to its possible candidacy as a magnetic refrigerant material.^{9–13} Stoichiometric Ni₂MnGa has an $L2_1$ crystal structure at room temperature and, upon cooling, it undergoes a first order martensitic structural phase transition (at T_M =202 K) from the parent cubic (austenitic) phase to a low temperature (LT) complex tetragonal structure. The Curie temperature for this is alloy is $T_c=376$ K.¹⁴ The substitution of Ni for Mn in the Ni_{2+x}Mn_{1-x}Ga system results in an increase in T_M and decrease of T_C and, for $0.17 \le x \le 0.20$, the coincidence of T_M and T_C results in a first order magnetostructural phase transition.¹⁵ As a result of this first order transition, a giant MCE of $|\Delta S_m| = 15 \text{ J kg}^{-1} \text{ K}^{-1}$ at 1.8 T field was observed in Ni_{2 19}Mn_{0.81}Ga.¹⁰ In stoichiometric $Ni_{55,2}Mn_{18,6}Ga_{18,6}$ a magnetic entropy change of $\Delta S_m =$ -20.4 J kg⁻¹ K⁻¹ at 317 K in a field change of 5 T has been reported.9 In polycrystalline Ni-Mn-Ga, the highest value of $\Delta S_m = -66.2 \text{ J kg}^{-1} \text{ K}^{-1}$ at 350.25 K in a field change of 5 T is observed in Ni_{2.19}Mn_{0.81}Ga.¹³ The coincidence of T_M and T_C is also reported to be the result of Cu substitution on the Mn sites of Ni₂MnGa.¹⁶ In a recent study, a giant MCE of $\Delta S_m \approx -64 \text{ J kg}^{-1} \text{ K}^{-1}$ at 308 K was observed in Ni₂Mn_{0.75}Cu_{0.25}Ga.¹⁷ Since the giant MCE occurs very close to room temperature, further research on and development of this alloy might result in the outcome of a potential magnetic refrigerant material that would be affordable and efficient. The ability to tune the magnetostructural transition temperature while preserving the high MCE value would be an interesting and significant outcome of further research. This is because the tunability of the high MCE value over a wide temperature range will open possibilities of developing magnetic refrigerant composites for near room temperature magnetic cooling applications.

In this work we report our experimental results of MCE studies on the Fe doped system (Ni₂Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga) and on a Ge doped system (Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}). The objective was to be able to tune the first order magnetostructural transition temperature through various substitutions while preserving the high ΔS_M peak value.

Polycrystalline buttons of approximately 5 g of $Ni_2Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga$ and $Ni_2Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}$ were fabricated by conventional arc melting in an argon atmosphere using Ni, Mn, Cu, Fe, Ga, and Ge of 4*N* purity. The elements were melted four times, and the weight loss after melting was found to be less than 0.3%. For homogenization, the samples were wrapped in Ta foil and annealed in vacuum for 72 h at 800 °C, and subsequently slowly cooled down to room temperature.

For phase identification and lattice constants determination, x-ray diffraction measurements were conducted at room temperature using a GBC minimaterials analyzer (MMA) x-ray diffractometer that employed Cu $K\alpha$ radiation and Bragg-Brentano geometry.

The magnetization measurements were performed using a superconducting quantum interference device (SQUID) made by Quantum Design, Inc. The measurements were performed in a temperature range of 5-400 K and magnetic

^{a)}Author to whom correspondence should be addressed; electronic mail: mkhan@siu.edu

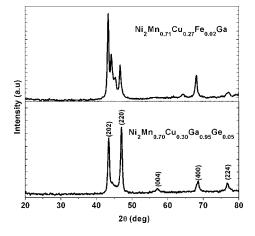


FIG. 1. Room temperature powder XRD patterns of $Ni_2Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga$ and $Ni_2Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}$.

field of up to 5 T. The magnetic entropy change ΔS_m was calculated from the isothermal magnetization data using the relation,

$$\Delta S_{\text{mag}} = \int_{0}^{H} \left(\frac{\partial M}{\partial T}\right)_{H} dH.$$
 (1)

Thermal expansion measurements were performed using a high resolution capacitance dilatometry method in the temperature range of 150–350 K.

Figure 1 represents the room temperature x-ray diffraction (XRD) patterns of Ni₂Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga and Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}. The XRD patterns indicate that Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05} possess a typical tetragonal structure at room temperature with a=b=5.46 Å and c= 6.41 Å. In the Ni₂Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga sample, both martenstic and austenitic phases seem to coexist. This coexistence of both phases is due to the martensitic transition taking place at T_M =302 K, which is very close to room temperature. A similar XRD pattern was observed for Ni₂Mn_{0.75}Cu_{0.25}Ga.¹⁶

The magnetization curves as a function of temperature M(T) of Ni₂Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga, Ni₂Mn_{0.75}Cu_{0.25}Ga and Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05} at a field of 1 kOe are presented in Fig. 2. As shown in this figure, the only transition observed in each alloy is a sharp jump of magnetization at T_c . The temperature hysteresis observed in the M(T) curves at 1 kOe, obtained for increasing and decreasing temperature, of Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05} [see inset (a) of Fig. 2] suggests that the transition at T_C is a first order phase transition. The sharp steplike change in the thermal expansion curve (typical for a first order phase transition) of $Ni_2Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}$ shown in inset (b) of Fig. 2 further verifies that the transition is a first order phase transition. It is clear from the figure that the magnetostructural transition temperature, represented by T_C , increases in the Ge doped sample and decreases in the Fe doped sample. It has been reported previously that Fe substitution on the Mn sites of Ni₂MnGa results in decrease of T_M and increase of T_C .¹⁸ In Ni₂Mn_{0.75}Cu_{0.25}Ga, $T_M = T_C$, and therefore as suggested in Ref. 16, the further partial substitution of Cu on the Mn sites results in the increase of T_M and decreases of T_C . Additional

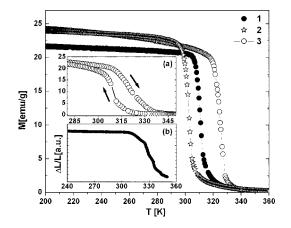


FIG. 2. Magnetization function of as а temperature of Ni₂Mn_{0.75}Cu_{0.25}Ga, (2) Ni₂Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga, (1)and (3)Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05} obtained at a field of 1 kOe. The inset (a) and (b) represents the temperature hysteresis and thermal expansion curves, respectively, of Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}.

substitution of Fe on the Mn site increases the T_C and decreases of T_M , bringing back the two transitions at the same temperature. Ge substitution on the Ga site in Ni₂MnGa results in a decrease of T_M ,¹⁹ and so when Ga is replaced by partial Ge in Ni₂Mn_{0.70}Cu_{0.30}Ga, T_M decreases resulting in the overlap of T_M and T_C . The saturation moments of the samples at 5 K are 3.32, 3.27, and $3.16\mu_B/f.u.$ for Ni₂Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga, Ni₂Mn_{0.75}Cu_{0.25}Ga, and Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}, respectively. It is apparent that the sample with Fe possesses a higher saturation moment of the Ni₂Mn_{0.75}Cu_{0.25}Ga system can be increased by Fe doping.

The ΔS_m values were evaluated from isothermal magnetization curves using Eq. (1). This equation is more appropriate to calculate MCE in the vicinity of a second order phase transition. However, its employment in calculating ΔS_m in the vicinity of first order phase transitions is very common which, according to Gschneidner et al., is justified in cases where problematic discontinuities are not present in the phase transition.³ The majority of the reported ΔS_m values of Ni-Mn-Ga, and other ferromagnetic systems³ exhibiting first order phase transitions, are calculated using Eq. (1). Figures 3(a) and 3(b) show the isothermal magnetization curves as a function of fields of Ni₂Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga and Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}, respectively. The changes entropies of Ni₂Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga, of magnetic Ni₂Mn_{0.75}Cu_{0.25}Ga, and Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}, as a function of temperature, are shown in Fig. 4. As shown in this figure, all of the samples possess comparable peak values of ΔS_m . Peak values of 32 and 58 J/kg K for Ni₂Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga and 23 and 57 J/kg K for Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05} are obtained at 2 and 5 T fields, respectively. The values compare well to those of Ni₂Mn_{0.75}Cu_{0.25}Ga (28 and 64 J/kg K at 2 and 5 T fields, respectively). As shown in Fig. 5, the ΔS_m peak values of Ni₂Mn_{0.75}Cu_{0.25}Ga and Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05} are found to be linearly dependent on the applied fields, whereas the ΔS_m peak values of Ni₂Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga are not linear with field. Due to this, at 2 T field the ΔS_m peak value of

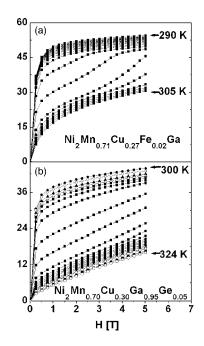


FIG. 3. Isothermal magnetization curves of (a) $Ni_2Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga$ and (b) $Ni_2Mn_{0.70}Cu_{0.30}Ga_{0.05}Ge_{0.05}$ at temperature increments of 1 and 0.5 K.

 $Ni_2Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga$ is observed to be larger than those of $Ni_2Mn_{0.75}Cu_{0.25}Ga$ and $Ni_2Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}$. This could be attributed to the nonlinear isothermal magnetization curves near the magnetostructural transition temperature of $Ni_2Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga$. The isothermal magnetization curves near magnetostructural transition temperature of $Ni_2Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}$ are found to be very linear.

We have studied magnetocaloric effects in $Ni_2Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga$ and $Ni_2Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}$. The manipulation of the Mn–Cu subsystem of Ni₂Mn_{0.75}Cu_{0.25}Ga with Fe doping results in a decrease of the first order magnetostructural transition temperature, whereas the substitution of Ge in the Mn-Cu-Ga subsystems results in an increase of the magnetostructural transition temperature. In both cases the giant magnetocaloric effect is successfully preserved. These experimental results suggest the possibility of tuning the first order magnetostructural transition temperatures while preserving the high MCE values in Ni₂MnGa Heulser alloys. We believe that these

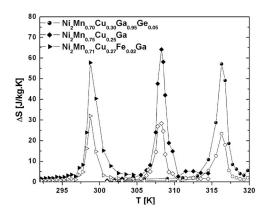


FIG. 4. Magnetic entropy changes (ΔS_M) as a function of temperatures of Ni₂Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga and Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05} for a field changes (ΔH) of 5 T (closed symbols) and 2 T (open symbols).

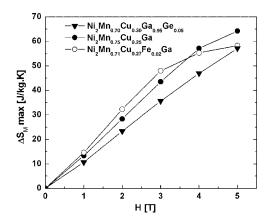


FIG. 5. Maximum magnetic entropy changes (ΔS_M max) as a function of fields of Ni₂Mn_{0.71}Cu_{0.27}Fe_{0.02}Ga and Ni₂Mn_{0.70}Cu_{0.30}Ga_{0.95}Ge_{0.05}.

results will significantly contribute to the understanding of the fundamental phenomenon of the phase transitions and related MCE in Ni–Mn–Ga based Heusler alloys, and thus will facilitate the development of promising magnetic refrigerants for near room temperature magnetic refrigeration applications.

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