Taming the First-Order Transition in Giant Magnetocaloric Materials

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The change in temperature resulting from a change in the magnetic field applied to a magnetic material is called the magnetocaloric effect (MCE). Integrating this effect into a thermodynamic cycle could lead to a broad range of applications. One of the most attractive is magnetic refrigeration at room temperature. This cooling technique offers the possibility of advantageously replacing the conventional vapor-compression technology since it has a higher efficiency and does not use refrigerant gases that are greenhouse gases.\(^1\)\(^2\) The search for magnetic materials displaying a large MCE is the first step in its development and is nowadays very active. More precisely, since the discovery of “giant” MCE (G-MCE), which occurs when the latent heat of a first-order transition (FOT) is involved,\(^3\) research has focused on reaching even stronger magnetic FOTs, that is to say, on reaching the largest latent heat possible, and the strongest magneto-structural coupling (e.g., cell-volume change at the FOT).\(^4\) However, though this approach is beneficial for reaching high MCE performances, in particular large entropy-change (\(\Delta S\)) values, an important drawback is the negative role of hysteresis. Since all MCE applications have a cyclic character, one of the main pre-conditions is to ensure a total (or at least partial) reversibility of the effect when either field or temperature oscillations are applied. From a material point of view, this means keeping the field or thermal hysteresis that could occur as small as possible. A second drawback of G-MCE materials is related to their mechanical stability. FOTs bring not only sharp magnetization jumps but also discontinuities of other physical parameters, including the unit cell. This “structural” part can have manifold aspects: symmetry breaking or cell-volume or lattice-parameter changes. The most dramatic for the stability of polycrystalline bulk samples turns out to be the cell-volume change. During thermal or magnetic field cycles, the strains generated by a volume change may cause fractures or even destruction of the bulk piece, which severely hinders the applicability of these materials. Technical solutions can be used to overcome this problem, for instance by embedding the MCE material in a resin or by a porous shaping.\(^19\) However, in such cases the MCE is “diluted”, which is not satisfactory since the gap of the magnet is not efficiently used and the thermal conductivity governing the heat transfer is decreased. Bulk G-MCE materials with a good mechanical stability should remain the preferred solution. Finally, to allow large-scale applications, a last requirement that should be borne in mind is that the MCE material must consist of elements that are available in large amounts, are not expensive, and are not classified as toxic.

In this context, the MnFe(P,x) system appears to be an ideal playground. This material family is derived from the Fe\(_7\)P compound, a prototypical example known for a long time to exhibit a sharp but weak (the latent heat \(L\) is only 0.25 \(kJ \cdot kg^{-1}\)) FOT with a Curie temperature (\(T_C\)) of 217 K.\(^19\) In this hexagonal system, the Fe atoms occupy two inequivalent atomic positions referred to as 3f (in a tetrahedral environment of non-metallic atoms) and 3g (pyramidal). An intriguing aspect is the disappearance of the magnetic moments of the iron atoms at the 3f sites, whereas there is only a limited decrease of the moments at the 3g site. This theoretical prediction has led to a cooperative description of the FOT that links the loss of long-range magnetic order at \(T_C\) with the loss of the local moments at the 3f site.\(^20\) This mechanism has recently been proposed to be the origin of the G-MCE observed in MnFe(P,Si). The disappearance of the magnetic moments is ascribed to a conversion from non-bonding d electrons to a distribution with pronounced hybridization with the surrounding Si/P atoms.\(^11\) A practical consequence is that the FOT mechanism can be expected to be highly sensitive to substitutions at the non-metallic site. In the present work, precisely this approach has been used to solve three problems of the previous generation of MnFe(P,Si) materials. 1) The first improvement needed is...
related to the enhancement of the adiabatic temperature change ($\Delta T_{ad}$). For magnetic-refrigeration applications, it is of primary interest to optimize the heart of the device, the MCE material. This MCE is quantified either by the entropy change ($\Delta S$) or by $\Delta T_{ad}$, depending on whether the magnetic-field change is performed isothermally or adiabatically. Comparing different materials reveals that a large $\Delta S$ does not necessarily imply a large $\Delta T_{ad}$ and vice versa; both quantities have to be optimized simultaneously. This is an unsatisfactory point for MnFe(P,Si) materials, which exhibit large $\Delta S$ but limited $\Delta T_{ad}$ (~2 K in 1 T).

2) The second target is to solve the mechanical stability problem observed in MnFe(P,Si). Even though the cell-volume change of these materials at the FOT is limited ($\Delta V = +0.2\%$), $\Delta T_{ad}$ crossing the transition still leads to fractures or destruction of bulk samples.

3) The last objective is to provide materials having a large MCE at intermediate fields, that is, at those offered by permanent magnets. If the magnetic field required to fully induce the FOT is considerably higher than the field source of the magnetic refrigerator (which is often the case for a FOT with large $L$), the material is designed in an inefficient way, since all the drawbacks of a FOT are present but only a part of the transition is really used. A good strategy to incorporate most of the latent heat into the MCE is to optimize the temperature shift of the transition due to the applied field.

Based on the Clausius–Clapeyron description $dT_C/dB = -T_C\Delta M/L$, where $B$ is the magnetic flux density and $\Delta M$ the jump in magnetization, the optimizations (1) and (3) above imply that $dT_C/dB$ should be increased, that is, an increase of $\Delta M$ and decrease of the latent heat. In earlier studies of MnFe(P,Si) materials, reduction of the hysteresis was achieved by adjustment of the Mn/Fe and P/Si ratios, which, unfortunately, reduces the magnetization. To increase the magnetization and $\Delta M$, it is required to work around the MnFeP$_{2/3}$Si$_{1/3}$ composition, which is a double optimum for the magnetization unmodified. For this purpose, boron substitution at the non-magnetic site (replacement of P by B) has been investigated.

In MnFe(P,Si) materials, the effect of B substitution on $T_C$ similar to that in Fe$_3$P.[23] On the isofield $M_B(T)$ curves, Figure 1, starting from the “magnetization optimum” MnFe$_{0.95}$P$_{0.67}$Si$_{0.33}$, substitution of B for 0.075 P (MnFe$_{0.95}$P$_{0.595}$B$_{0.075}$Si$_{0.33}$) leads to a significant increase of $T_C$ (for $T_C$ measured upon heating from 215 K to 283 K; see the Supporting Information S1 for intermediate compositions). Most interesting is the fact that the extremely large difference between heating and cooling Curie temperatures of the parent compound MnFe$_{0.95}$P$_{2/3}$Si$_{1/3}$ (thermal hysteresis of 75 K) is strongly suppressed by B substitution, resulting in a hysteresis of only 1.6 K in the B = 7.5% compound. The average hysteresis decrease by B doping is about 10 K per percent B, a value much higher than obtained by changing the Mn/Fe ratio (~1.2 K per Mn or Fe%) or the P/Si ratio (~1.1 K per Si%). Moreover, since the control of the hysteresis is now achieved without deviating from the Mn/Fe ~ 1 and Si ~ 1/3 optima, the magnetization in the ferromagnetic state far below $T_C$ is kept unmodified (the two samples exhibit a magnetization of 150 A m$^{-2}$ kg$^{-1}$ at 50 K), which is very important for the MCE performance.

The MCE evaluation of a new material requires a complete characterization of both $\Delta S$ and $\Delta T_{ad}$. Figure 2 combines results from two direct probes and from indirect (magnetic and calorimetric (differential scanning calorimetry, DSC)) techniques. Such an extensive characterization has only recently been published for G-MCE materials,[24] but is crucial in order to obtain a reliable picture of the thermomagnetic behavior around a FOT. The specific-heat curves of MnFe$_{0.95}$P$_{0.595}$B$_{0.075}$Si$_{0.33}$ can be found in the Supporting Information. The latent heat obtained by the integration of the curve in zero field is $L = 3.8$ kJ kg$^{-1}$. This significantly lower value than for materials without B ($L = 12.3$ kJ kg$^{-1}$ for MnFe$_{0.95}$P$_{2/3}$Si$_{1/3}$ and $L = 6.4$ kJ kg$^{-1}$ for Mn$_{1.25}$Fe$_{0.75}$P$_{1/2}$Si$_{1/2}$) is assumed to be at least partially the reason for the larger $dT_C/dB = 4.3(2)$ K T$^{-1}$ found in MnFe$_{0.95}$P$_{0.595}$B$_{0.075}$Si$_{0.33}$ than the $3.5(2)$ K T$^{-1}$ found in Mn$_{1.25}$Fe$_{0.75}$P$_{1/2}$Si$_{1/2}$.[23] From a metrological point of view, special care has been paid to avoid miscalculations that could result in a “spike” on the $\Delta S(T)$ curves.[25–27] Moreover, since both the hysteresis and the width of the FOT are limited in the present compound, one can expect this problem to be very minor.

The $\Delta S(T)$ curves derived from isofield magnetic measurements and calorimetry upon heating, Figure 2a, are in good agreement: $\Delta S_{max} = 9.1(8)$ J kg$^{-1}$ K$^{-1}$ at 279.1 K by DSC and $\Delta S_{max} = 9.8 \pm 1.0$ J kg$^{-1}$ K$^{-1}$ at 281 K by the magnetic method. These values are much higher than those of the archetypical MCE material Gd.[1,2] Compared with G-MCE materials, $\Delta S$ of MnFe(P,Si,B) is similar to that of Gd$_3$Si$_2$Ge$_2$, higher than the values generally found in Heusler alloys, and just below the best performances found for La(Fe,Si)$_3$H$_6$.[1,2] However, it should be kept in mind that the indirect $\Delta S$ measurements we are referring to do not disclose the reversibility of the MCE, a point for which direct measurements, in particular of $\Delta T_{ad}$, are more suitable.

We used two types of direct methods to measure the adiabatic temperature change (denoted as $\Delta T_{cyclic}$ hereafter). The first direct measurements were carried out in a device using a

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**Figure 1.** Temperature dependence of the magnetization in the parent compound and in a boron-substituted material. Isofield magnetization curves recorded at $B = 1$ T upon cooling (filled symbols) and upon heating (open symbols) for MnFe$_{0.95}$P$_{0.67}$Si$_{0.33}$ (squares) and MnFe$_{0.95}$P$_{0.595}$B$_{0.075}$Si$_{0.33}$ (triangles).
continuous sweep of the temperature during field oscillations \( (\Delta B = 1.1 \, \text{T}) \). Figure 3a presents a typical raw-data curve. Application and removal of the magnetic field creates a tower shape response overlaying a linear temperature versus time evolution, which corresponds to the sweep rate of the measurement. The same values \( \Delta T_{\text{cyclic}} = 2.8 \, \text{K} \) are obtained irrespectively of the direction of the magnetic-field change, which is a first indication of total reversibility of the MCE in this material. To avoid any inconsistencies that may result from the dynamic character of this measurement, especially when dealing with a material still displaying a small thermal hysteresis, a second kind of this measurement, especially when dealing with a material of G-MCE compounds. 

For instance, the present performance are close to the "giant \( \Delta T_{\text{ad}} \)" reported in Heusler alloys \( (\Delta T_{\text{ad}} = 2.8 \, \text{K} \) at 1 T for \( \text{Ni}_{5.5}\text{Mn}_{26}\text{In}_{13}\text{Co}_{5} \)). However, in sharp contrast with this latter example, where only a small part of \( \Delta T_{\text{ad}} \) is reversible, the large MCE presently observed is fully reversible when the field is cycled, which is an overwhelming advantage of MnFe(P,Si,B) compounds.

These new materials are also a significant improvement over the previous MnFe(P,Si) generation, since the same \( \Delta S \) values are observed while \( \Delta T_{\text{ad}} \)
has been enhanced by more than 30%. This enhancement of ΔTad at intermediate magnetic fields is ascribed to the larger dTc/dB, so indirectly comes from the higher magnetization and lower latent heat of this new material.

The last progress achieved for MnFe(P,Si,B) materials concerns the mechanical stability. This problem, already noticed for G-MCE materials, manifests itself by, for instance, the degradation of ΔTad during cyclic measurements.[18] Qualitatively, we observe that bulk pieces of MnFe0.95P0.595B0.075Si0.33 do not show any degradation when cycled across the transition. Quantitatively, even after 10000 cycles of magnetization/demagnetization used in the ΔTcyclic measurements, the sample geometry remains intact and the mechanical integrity is maintained while the MCE does not show any degradation (Supporting Information S3). We believe that this improved mechanical stability in comparison with MnFe(P,Si) compounds or other G-MCE materials is partially related to the absence of a cell-volume change at the transition. In MnFe0.95P0.595B0.075Si0.33, the jump of the c/a ratio at Tc is almost as pronounced as in Mn1.25Fe0.7P0.5Si0.5,[21] but no discontinuous volume change was observed within the experimental accuracy, Figure 4. This exceptionally small ΔV turns out to be smaller than in all other MnFe(P,x) compounds, for instance smaller than in As-based materials (where ΔV/V = −0.44%),[29] in Ge-based materials (where ΔV/V = +0.1%),[20] and in Si-based materials.[21]

Generally speaking, the present study highlights that, besides magnetism, the key parameter to be controlled in the design of promising G-MCE materials is the latent heat of the FOT. More precisely, L has to be large enough to ensure high ΔS values but should be “kept under control” to maintain a large dTc/dB and ΔTad. Regarding the MnFe(P,x), MCE family, it is found that B substitution has a similar effect as in Fe2P, a point which was not straightforward at first glance since we are actually dealing with FOTs about 50 times stronger than in Fe2P. From an application point of view, the combination of a tunable Tc (Mn/Fe, P/Si, and B composition parameters), no toxic or critical elements, high ΔS, high ΔTad total reversibility of the effect, and good mechanical stability gives MnFe(P,Si,B) compounds considerable advantages, making them highly promising for magnetic refrigeration. Compared to La(Fe,Si)13 hydrides, it is still unclear which material has the largest reversible MCE but, in any case, these new MnFe(P,Si,B) materials neither contain a rare-earth element nor suffer from dehydrogenation.

Experimental Section

The parent and B-substituted samples were prepared in the same way. Stoichiometric quantities of Mn flakes, Fe2P, B, and Si powders were ground in a planetary ball mill for 10 h. The resulting powders were pressed into pellets and sealed in quartz ampules under 200 mbar of Ar. Double-step sintering was performed according to the process already described.[21] The samples were cycled 5 times across the FOT prior to any measurements. The structural parameters were studied by collecting X-ray diffraction patterns at various temperatures in a PANalytical X’Pert PRO diffractometer equipped with an Anton Paar TTK450 low-temperature chamber. The magnetization measurements were carried out in a magnetometer equipped with a superconducting quantum interference device (SQUID; Quantum Design MPMS 5 XL with reciprocating sample option, RSO). The isofield magnetization curves Mx(T) were measured in sweeping mode at a rate of 1 K min−1 with a temperature increment of 0.5 K. For the ΔS derivation on the basis of magnetic data and the use of the Maxwell equation,[2] Mx(T) curves were recorded with a field increment of 0.25 T. The DSC measurements were performed in a Peltier cell calorimeter described previously,[29] from which calorimetric ΔS and ΔTad were derived by means of the usual methods.[2] The direct ΔTcyclic measurements with temperature sweeping, Figure 3a, were carried out in a homemade device, for which the temperature versus time signal was recorded by a thermocouple clamped between two slices of sample (cylindrical shape of 10 mm diameter and 1 mm thickness). External-magnetic-field changes were applied by moving/removing the samples from the magnetic field at a rate of 1.1 T s−1. The direct ΔTcyclic measurements of Figure 3b were carried out on a cubic sample of 60 mg with the probe described in the literature.[31]

Supporting Information

Supporting Information is available from the Wiley Online Library or from the author.

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