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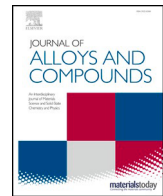
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# Ultra-low hysteresis in giant magnetocaloric $Mn_{1-x}V_xFe_{0.95}(P,Si,B)$ compounds



Jiawei Lai<sup>a,b</sup>, Xinmin You<sup>b</sup>, Jiayan Law<sup>c</sup>, Victorino Franco<sup>c</sup>, Bowei Huang<sup>a,b</sup>,  
Dimitrios Bessas<sup>d</sup>, Michael Maschek<sup>b</sup>, Dechang Zeng<sup>a,\*</sup>, Niels van Dijk<sup>b</sup>, Ekkes Brück<sup>b,\*</sup>

<sup>a</sup> School of Materials Science & Engineering, South China University of Technology, Guangzhou 510640, China

<sup>b</sup> Fundamental Aspects of Materials and Energy, Department of Radiation Science and Technology, TU Delft, Mekelweg 15, 2629JB Delft, the Netherlands

<sup>c</sup> Dpto. Física de la Materia Condensada ICMSE-CSIC, Universidad de Sevilla, Apdo1065, 41080 Sevilla, Spain

<sup>d</sup> European Synchrotron Radiation Facility, F-38043 Grenoble, France

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## ABSTRACT

Large thermal hysteresis in the (Mn,Fe)<sub>2</sub>(P,Si) system hinders an efficient heat exchange and thus limits the magnetocaloric applications. Substitution of manganese by vanadium in the  $Mn_{1-x}V_xFe_{0.95}P_{0.593}Si_{0.33}B_{0.077}$  and  $Mn_{1-x_2}V_{x_2}Fe_{0.95}P_{0.563}Si_{0.36}B_{0.077}$  compounds enable a significant reduction in the thermal hysteresis without losing the giant magnetocaloric effect. For the composition closest to the critical one, where first-order crossovers to second-order phase transition in the series of  $x_2 = 0.02$ ,  $Mn_{0.98}V_{0.02}Fe_{0.95}P_{0.563}Si_{0.36}B_{0.077}$  exhibits a thermal hysteresis that is reduced from 1.5 to 0.5 K by 67%, yielding an adiabatic temperature change of 2.3 K and magnetic entropy change of 5.6 J/kgK for an applied field of 1 T, which demonstrates its potential for highly efficient magnetic heat pumps utilizing low-cost permanent magnets.

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## 1. Introduction

Room temperature magnetic heat pumps attract broad attention due to their advantages compared to the traditional heat pumps, such as a high efficiency and a low impact on the environment [1,2]. Materials with a giant magnetocaloric effect (GMCE), which are the working materials of this technique, will release heat when an external magnetic field is applied, while they absorb heat when the magnetic field is removed. The performance of GMCE materials mainly determines the working efficiency and the cooling power of this technology. The GMCE usually occurs in materials that show a first-order magnetic transition (FOMT), such as  $Gd_5Ge_2Si_2$ , [3]  $LaFe_{13-x}Si_x$ , [4–7]  $MnFeP_{1-x}As_x$ , [8]  $MnFeP_{1-x-y}Si_xB_y$ , [9–11]  $MnCoGe$  ( $Cu, B$ )<sub>x</sub> [12,13] and Heusler [14] compounds. Among them, the  $MnFeP_{1-x-y}Si_xB_y$  compounds are regarded as one of the most promising materials that can be industrialized because of their cheap and non-toxic elements, high cooling capacity and tunable  $T_c$  near room temperature [9].

$MnFeP_{1-x}Si_x$  compounds with a strong FOMT show a significant change in lattice parameters across the magnetic phase transition

while keeping its hexagonal structure, which is a magneto-elastic transition [15]. However, a strong FOMT is often accompanied by large hysteresis, including both magnetic field hysteresis and thermal hysteresis ( $\Delta T_{hys}$ ), which will reduce field-induced temperature change. Hysteresis limits the practical application of these materials since it will largely reduce the working efficiency [16]. For some first-order materials that undergo a first-order magneto-structural transition, for instance Heusler alloys, the thermal hysteresis and magnetic field hysteresis may show a very different behavior. The magneto-structural transition is suppressed in field and thus a low hysteresis at low applied field and a larger hysteresis at high applied field could occur [17]. However, the presently studied (Mn,Fe)<sub>2</sub>(P,Si) based alloys undergo a magneto-elastic transition without a structural transition. In magneto-elastic transitions the thermal hysteresis and the magnetic field hysteresis are generally closely coupled, as reported previously [18]. We therefore consider that the low thermal hysteresis  $\Delta T_{hys}$  is representative for a low magnetic field hysteresis, and thereby provides important information for the thermodynamic properties for magnetic heat pumps and magnetic refrigerators.

Theoretically, for a thermal hysteresis  $\Delta T_{hys}$  of 2 K, the efficiency can be 60% of the Carnot efficiency and it can reach 75% of the Carnot efficiency if a value of 0.5 K is achieved [16]. Therefore, in the premise of keeping the GMCE, the thermal hysteresis should be made as narrow as possible by tailoring the microstructure or by tuning the

\* Corresponding authors.

E-mail addresses: [medczeng@scut.edu.cn](mailto:medczeng@scut.edu.cn) (D. Zeng),  
[E.H.Bruck@tudelft.nl](mailto:E.H.Bruck@tudelft.nl) (E. Brück).

chemical composition. Recently, through 0.075 at% of B substitution in the  $\text{MnFe}_{0.95}\text{P}_{0.595}\text{Si}_{0.33}\text{B}_{0.075}$  compounds,  $\Delta T_{\text{hys}}$  can be optimized to 1.6 K according to the temperature-dependent magnetization curves and to 2.0 K according to in-field DSC measurements, both at magnetic field of 1 T (see the supporting information of reference [11]). This compound is generally regarded as one of the benchmarks for the  $\text{Fe}_2\text{P}$ -type compounds. It can be cycled for 10 thousand times and the sample geometry remains intact. A higher level of B substitution can further decrease  $\Delta T_{\text{hys}}$  but fails to provide a sufficiently large GMCE [17]. It is essential to find a new approach to further decrease  $\Delta T_{\text{hys}}$  and simultaneously provide a sufficiently high GMCE, as characterized by the isothermal magnetic entropy change ( $\Delta S_M$ ) and the adiabatic temperature change ( $\Delta T_{\text{ad}}$ ). To evaluate the size of  $\Delta S_M$ , Gd with a value of  $\Delta S_M$  is 3 J/kgK under 1 T is generally used as a standard to compare with [18]. Another design criterion is that  $\Delta T_{\text{ad}}$  should be larger than 2.0 K [19], since Engelbrecht and Bahl [20] demonstrated that cooling (and heating) is ineffective when  $\Delta T_{\text{ad}}$  drops below 2.0 K.

On the other hand, the magnetic field currently used in the prototypes of heat pumps are generated by NdFeB permanent magnets, which varies from 0.9 to 1.9 T [21–24]. The overall cost to reach a field of 1.9 T can be 10 times higher than the cost to reach a field of 0.9 T. It is therefore of interest to explore the lower magnetic field potential of the magnetic heat pump system.  $dT_c/dB$  is defined as the shift of  $T_c$  induced by an applied magnetic field. As the magnitude of  $dT_c/dB$  is regarded as the driving force of the magneto-elastic transition [25], if enhanced, the magnetic phase transition can be induced in lower magnetic fields. It is positive for conventional first-order magnetic transition materials, such as  $\text{MnFeP}_{1-x}\text{ySi}_x\text{By}$  [11] and La-Fe-Si [26], while it is negative for the inverse first-order magnetic transition materials, for instance the Ni–Mn–X–Heusler compounds with X = Sn, Sb and In [27] or Fe–Rh [28]. For the conventional first-order magnetic transition materials, this shift is attributed to the magnetic field stabilization of the phase with the higher magnetization, being the low-temperature ferromagnetic phase [13,29]. In an external magnetic field, extra thermal energy is then needed to induce the magnetic phase transition. Thus,  $T_c$  shifts to higher temperature. One of our motivations of this work is to increase the driving force (i.e.  $dT_c/dB$ ) of FOMT and make it feasible to be utilized in devices with low-cost permanent magnets.

In previous reports we have shown an enhanced GMCE in the low magnetic field range by introducing V substitution to Mn-Fe-P-Si compounds [30,31]. From the motivation of minimizing  $\Delta T_{\text{hys}}$  and achieving a large GMCE under a magnetic field of 1.0 T, the combined effect of B and V additions to the Mn-Fe-P-Si compounds is investigated in this study. The effect of V substitution on the thermal hysteresis  $\Delta T_{\text{hys}}$ , the lattice parameters, the magnetic properties and  $dT_c/dB$  in polycrystalline Mn-V-Fe-P-Si-B compounds are systematically investigated. Through optimization of the composition and V substitution, an ultra-low  $\Delta T_{\text{hys}}$  of 0.5 K and a large GMCE with  $|\Delta S_M| = 6.2$  J/kgK and  $\Delta T_{\text{ad}} = 2.5$  K at an internal magnetic field of 1 T is achieved simultaneously.

## 2. Experimental

Polycrystalline  $\text{Mn}_{1-x_1}\text{V}_{x_1}\text{Fe}_{0.95}\text{P}_{0.593}\text{Si}_{0.33}\text{B}_{0.077}$  ( $x_1 = 0.00, 0.02, 0.03$ ) compounds were prepared by a powder metallurgy method. The vanadium substitution started from 0.02 since it was reported to be optimal to tune the hysteresis and entropy change in low magnetic fields [30]. The starting materials in the form of Mn, Fe, red P, Si and V powders were mechanically ball milled for 10 h in an Ar atmosphere with a constant rotation speed of 380 rpm, then pressed into small pellets, and finally sealed in quartz ampoules under 200 mbar of Ar before employing the various heat treatment conditions. The pellets were annealed at 1323 K for 2 h to promote crystallization and cooled to room temperature in the furnace. Then they

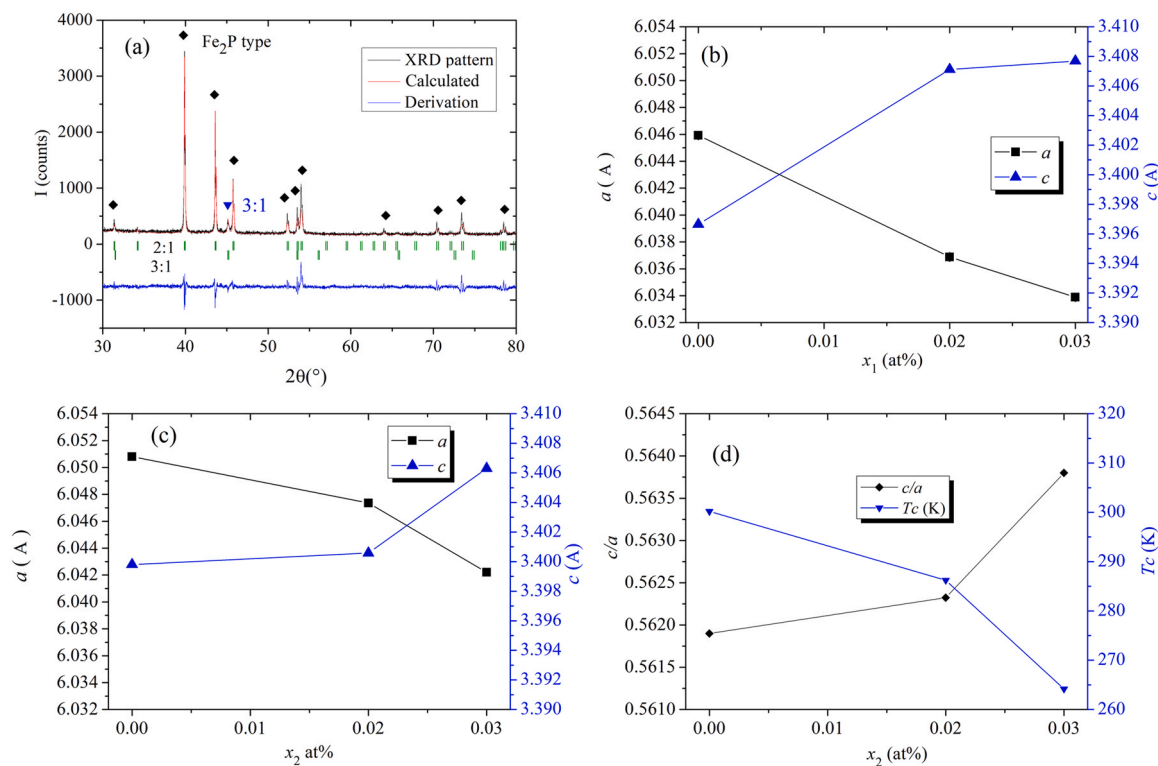
were heated up to the same annealing temperature for 20 h to homogenize the compound and finally quenched in water. This batch samples were labeled as *series A*. In this series, the composition of sample with  $x_1 = 0.00$  is designed to be the same as to the previous benchmark material  $\text{MnFe}_{0.95}\text{P}_{0.595}\text{Si}_{0.33}\text{B}_{0.075}$  [11]. In order to enhance the  $|\Delta S_M|$ , an optimal composition series of  $\text{Mn}_{1-x_2}\text{V}_{x_2}\text{Fe}_{0.95}\text{P}_{0.563}\text{Si}_{0.36}\text{B}_{0.077}$  ( $x_2 = 0.00, 0.02, 0.03$ ) compounds with a higher Si content was prepared with the same procedure as *series A* at a higher annealing temperature of 1373 K. This batch samples are labeled as *series B*.

X-ray diffraction (XRD) patterns were collected on a PANalytical X-pert Pro diffractometer with Cu-K $\alpha$  radiation (1.54056 Å) at room temperature. The temperature dependence and magnetic field dependence of the magnetization were measured with a commercial superconducting quantum interference device (SQUID) magnetometer (Quantum Design MPMS 5XL) in the reciprocating sample option (RSO) mode. The adiabatic temperature change ( $\Delta T_{\text{ad}}$ ) was measured in a Peltier cell based differential scanning calorimeter (DSC) using a Halbach cylinder providing a magnetic field of up to 1.5 T. In this setup, the iso-field calorimetric scans were performed at a rate of 3 K min $^{-1}$ , while the temperature had been corrected for the effect of the thermal resistance of the Peltier cells.

## 3. Results and discussion

Fig. 1(a) shows the room temperature XRD patterns and the refined results for the  $\text{Mn}_{1-x_2}\text{V}_{x_2}\text{Fe}_{0.95}\text{P}_{0.563}\text{Si}_{0.36}\text{B}_{0.077}$  ( $x_2 = 0.02$ ) compound. The main phase is identified as the hexagonal  $\text{Fe}_2\text{P}$ -type phase (2:1) with a space group of P-62 *m* and the impurity phase is the cubic  $\text{MnFe}_2\text{Si}$ -type phase (3:1) with a space group of *Fm*3*m*, which agrees with the previously reported data [1]. A high weight fraction of 96.3 wt% is obtained for the main phase, suggesting that the annealing condition and composition are suitable to achieve a high purity of  $\text{Fe}_2\text{P}$ -type phase. Based on the refinement results, the weight fraction of the impurity phase is  $1.8 \pm 0.1$ ,  $2.4 \pm 0.1$  and  $2.0 \pm 0.1$  wt% for  $x_1 = 0.00, 0.02$  and  $0.03$  and is  $4.5 \pm 0.1$ ,  $3.8 \pm 0.1$  and  $4.3 \pm 0.1$  wt% for  $x_2 = 0.00, 0.02$  and  $0.03$ , respectively. Thus, the variation in the amount of impurity phase by V substitution is negligible within the same series.

In Fig. 1(b), the results of the Fullprof refinement [32] for the lattice parameters of the *series A* compounds are shown. An increase in V substitution leads to a decrease in *a* axis and an increase in *c* axis. The lattice constant *a* decreases from 6.0459 to 6.0339 Å and *c* increases from 3.3966 to 3.4076 Å when  $x_1$  increases from 0 to 0.03. In Fig. 1(c) and 1(d), the lattice parameters, *c/a* ratio and  $T_c$  are shown for the *series B* compounds. The lattice constant *a* decreases from 6.0508 to 6.0422 Å and *c* increases from 3.3998 to 3.4063 Å when  $x_2$  is increased from 0 to 0.03, see the Fig. 1(c). The increase in the *c/a* ratio leads to a decrease in  $T_c$ , as shown in Fig. 1(d). The correlation between the *c/a* ratio and  $T_c$  originates from the distinct mixed magnetism of the  $\text{Fe}_2\text{P}$  type compounds [33]. The Fe atoms prefer to occupy the plane and Mn atoms prefer to occupy the 3*g* plane with P and Si randomly distributed in the hexagonal lattice. When transforming from the paramagnetic to the ferromagnetic state, the lattice constant *a* increases and causes a stronger localization for the 3*d* electrons of Fe (3*f*) instead of chemical bonding with non-metallic P and Si. Jointly with the lattice constant *c*, decreases the interlayer distance between the Mn and Fe atoms and thereby enhances the magnetic exchange interaction between two magnetic atoms [34]. Therefore, adding V to  $\text{Mn}_{1-x_2}\text{V}_{x_2}\text{Fe}_{0.95}\text{P}_{0.593}\text{Si}_{0.33}\text{B}_{0.077}$  compounds will increase the *c/a* ratio and leads to a reduction in the magnetic exchange interaction. Similar trend is observed for the  $\text{Mn}_{1-x_1}\text{V}_{x_1}\text{Fe}_{0.95}\text{P}_{0.593}\text{Si}_{0.33}\text{B}_{0.077}$  compounds. As a result,  $T_c$  is reduced. These trends are consistent with the previous investigation for the effect of V substitution in the Mn-Fe-P-Si compounds [30].

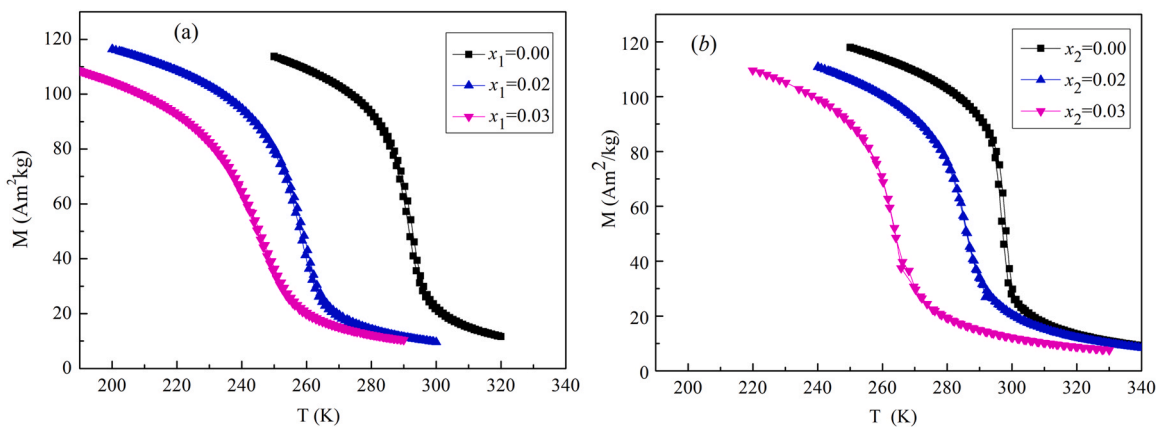


**Fig. 1.** (a) XRD pattern and the refined results for  $\text{Mn}_{1-x_2}\text{V}_{x_2}\text{Fe}_{0.95}\text{P}_{0.563}\text{Si}_{0.36}\text{B}_{0.077}$  ( $x_2 = 0.02$ ) compounds; (b) Lattice parameters  $a$  and  $c$  as function of value of  $x_1$  for  $\text{Mn}_{1-x_1}\text{V}_{x_1}\text{Fe}_{0.95}\text{P}_{0.593}\text{Si}_{0.33}\text{B}_{0.077}$  (series A) and (c) Lattice parameters  $a$  and  $c$  and (d)  $c/a$  ratio and  $T_C$  as function of value of  $x_2$  for  $\text{Mn}_{1-x_2}\text{V}_{x_2}\text{Fe}_{0.95}\text{P}_{0.563}\text{Si}_{0.36}\text{B}_{0.077}$  (series B).

The temperature dependence of the magnetization in the compounds of series A and B are shown in Fig. 2(a) and 2(b), respectively. For decreasing temperature, they have a temperature-induced magnetic phase transition from paramagnetic to ferromagnetic state. The transition temperature  $T_C$  is determined from the maximum value of  $-dM/dT$  in the  $M$ - $T$  curve measured during heating at 1 T. For series A,  $T_C$  tends to decrease with increasing V substitution. When  $x_1$  increases from 0 to 0.02,  $T_C$  decreases from 290.0 to 255.2 K. For series B,  $T_C$  decreases from 300 to 286.2 K when  $x_2$  increases from 0 to 0.02. Thus,  $T_C$  in series B is tuned to the near room-temperature region, which makes it suitable as a series of working materials for near room temperature applications. The value of  $-dM/dT$  indicates how strong the FOMT is. The values of  $-dM/dT$  are 7.6, 5.0 and 3.0  $\text{Am}^2/\text{kgK}$  for  $x_1 = 0.00, 0.02$  and 0.03 and are 11.1, 5.6 and

5.2  $\text{Am}^2/\text{kgK}$  for  $x_2 = 0.00, 0.02$  and 0.03, respectively. Thus,  $-dM/dT$  decreases by increasing V content, i.e. the FOMT is weakened.

The value of  $\Delta T_{\text{hys}}$  is determined by calculating the difference in the maximum value of  $-dM/dT$  during cooling and heating in an applied magnetic field of  $\mu_0 H = 1$  T at a ramp rate of 2  $\text{Kmin}^{-1}$ . For series A,  $\Delta T_{\text{hys}}$  decreases by 36% from 1.1 to 0.7 K when  $x_1$  increases from 0.00 to 0.03. For series B,  $\Delta T_{\text{hys}}$  decreases by 93% from 1.5 to 0.1 K when  $x_2$  increases from 0.00 to 0.03, see Table 1. The  $M$ - $H$  curves for series B are measured under ascending and descending fields around the Curie temperature using a loop process [35]. The magnetic hysteresis is also found to decrease when  $x_2$  increases from 0 to 0.02, as shown in the Fig. S1. Thus, magnetic and thermal hysteresis show the same trend. When a  $\Delta T_{\text{hys}}$  of 0.5 K is achieved, up to 75% of the Carnot efficiency is feasible for the sample with  $x_2 = 0.02$



**Fig. 2.** Temperature dependence of magnetization for (a)  $\text{Mn}_{1-x_1}\text{V}_{x_1}\text{Fe}_{0.95}\text{P}_{0.593}\text{Si}_{0.33}\text{B}_{0.077}$  and (b)  $\text{Mn}_{1-x_2}\text{V}_{x_2}\text{Fe}_{0.95}\text{P}_{0.563}\text{Si}_{0.36}\text{B}_{0.077}$  compounds under an applied magnetic field of 1 T.

**Table 1**

Curie temperature ( $T_C$ ), thermal hysteresis ( $\Delta T_{hys}$ ), latent heat ( $L$ ), magnetic entropy change ( $|\Delta S_M|$ ) and adiabatic temperature change  $\Delta T_{ad}$  at a field change of 1 T for series A and B. The value of  $\Delta T_{hys}$  is determined in an applied magnetic field of  $\mu_0 H = 1$  T at a ramp rate of 2 Kmin<sup>-1</sup>.

	Annealed T (K)	Sample	$T_C$ (K)	$\Delta T_{hys}$ (K)	$L$ (kJ/kg)	$ \Delta S_M $ (J/(kg·K))	$\Delta T_{ad}$ (K)
series A	1323	$x_1 = 0.00$	290.0	1.1	5.2	6.5	2.7
series A	1323	$x_1 = 0.02$	255.2	0.9	2.4	4.6	1.6
series A	1323	$x_1 = 0.03$	242.5	0.7	1.7	2.7	/
series B	1373	$x_2 = 0.00$	300.2	1.5	6.2	11.3	3.5
series B	1373	$x_2 = 0.02$	286.2	0.5	3.3	5.6	2.3
series B	1373	$x_2 = 0.03$	264.2	0.1	2.8	4.8	1.6
Ref. [11]	1373	$x_1 = 0.00$	281	1.6	3.8	9.8	2.5

[14]. The value of  $\Delta T_{hys}$  is determined by both intrinsic properties, i.e. the strength of the FOMT and extrinsic properties like the fraction of the impurity phase [13]. Here the intrinsic weakening of FOMT is the dominant factor that reduces  $\Delta T_{hys}$  to an ultra-low value as the compounds have very similar amount of impurity phase within the series.

The latent heat is an important quantity to characterize how strong the FOMT is. The DSC curves for series A and B are measured under zero field, as shown in the Fig. 3, and the derived latent heat is also listed in Table 1.

In Fig. 3, the DSC curves for series A and B were measured under zero field. By increasing the V content, the heat capacity is largely decreased, indicating that a strong FOMT is transferred to a weak FOMT. For the sample with  $x_1 = 0.00$  and  $x_2 = 0.00$ , sharp heat capacity peaks are shown, indicating a strong FOMT. When increasing V content, the heat capacity is decreased, suggesting the FOMT is weaker. As shown in the Table 1, increasing  $x_1$  from 0.00 to 0.03 results in a strong reduction of the latent heat by 67% from 5.2 to 1.7 J/g for series A. Similarly, increasing  $x_2$  from 0.00 to 0.03 results in a strong reduction of the latent heat by 55% from 6.2 to 2.8 J/g (listed in Table 1). The sample with  $x_2 = 0.02$  in series B has a slightly lower latent heat of 3.3 J/g than the previous benchmark MnFe<sub>0.95</sub>P<sub>0.595</sub>Si<sub>0.33</sub>B<sub>0.077</sub> compound of 3.8 J/g [11]. Therefore, the lower latent heat and the high fraction of the Fe<sub>2</sub>P-type phase (96.3 wt%) can explain the low value of  $\Delta T_{hys}$  of 0.5 K found in the sample with  $x_2 = 0.02$ .

The iso-thermal entropy change  $|\Delta S_M|$  is widely used as a criterion to evaluate the amount of heat transferred in one thermodynamic cycle. The values of  $|\Delta S_M|$  are derived from iso-field magnetization curves using the Maxwell relation [36,37]. The iso-field magnetization curves of series A and B for a magnetic field change are measured in the vicinity of  $T_C$  with a temperature interval

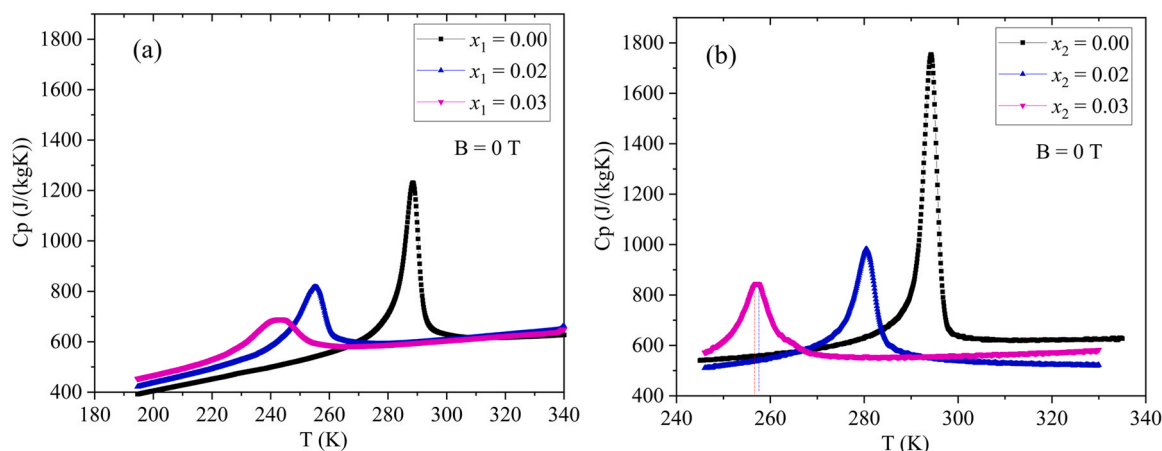
of 1 K. In order to minimize the effect of demagnetization factor, the applied magnetic field starts from 0.6 T and increases to 2 T in a step of 0.1 T. The calculated temperature dependence of  $|\Delta S_M|$  for series A and B are shown in Fig. 4(a) and 4(c), respectively. The values of  $|\Delta S_M|$  are 5.9, 4.5, 2.7 J/(kgK) at 1 T and 10.5, 8.0, 5.5 J/(kgK) at 2.0 T for samples with  $x_1 = 0, 0.02$ , and 0.03, respectively. Note that the GMCE of the compound with  $x_1 = 0.00$  is lower than the previous benchmark compound for Mn-Fe-P-Si-B [11] ( $|\Delta S_M| = 9.2$  J/(kgK) at 1 T), which is due to the decrease of annealing temperature from 1373 to 1323 K. In series B, values of  $|\Delta S_M|$  are 11.3, 5.6, 4.8 J/(kgK) at 1 T and 14.9, 8.4, 8.0 J/(kgK) at 2.0 T for samples with  $x_2 = 0, 0.02$ , and 0.03, respectively. After increasing the Si content from 0.33 to 0.36,  $|\Delta S_M|$  of the sample with  $x_2 = 0.00$  in series B outperforms the previously benchmark compound for Mn-Fe-P-Si-B [11]. This provides some room for V substitution to decrease  $\Delta T_{hys}$ , while retaining a large  $|\Delta S_M|$ . When  $x_2$  increases to 0.02,  $|\Delta S_M|$  is reduced to 5.6 J/(kgK) at 0–1 T, which is still about two times larger than that of polycrystalline Gd ( $|\Delta S_M| = 3.0$  J/(kgK) [18]).

In the magnetocaloric properties,  $\Delta T_{ad}$  illustrates the direct temperature span that can be achieved in the magnetic heat pump. In-field DSC measurements were conducted at a rate of 3 Kmin<sup>-1</sup> in the temperature range of 240–340 K. From these measurements,  $|\Delta S_M|$  and  $C_p$  were collected under a constant magnetic field ( $\mu_0 H = 0, 0.5, 1.0$  and 1.5 T). The values of  $\Delta T_{ad}$  were calculated by the following equation: [38].

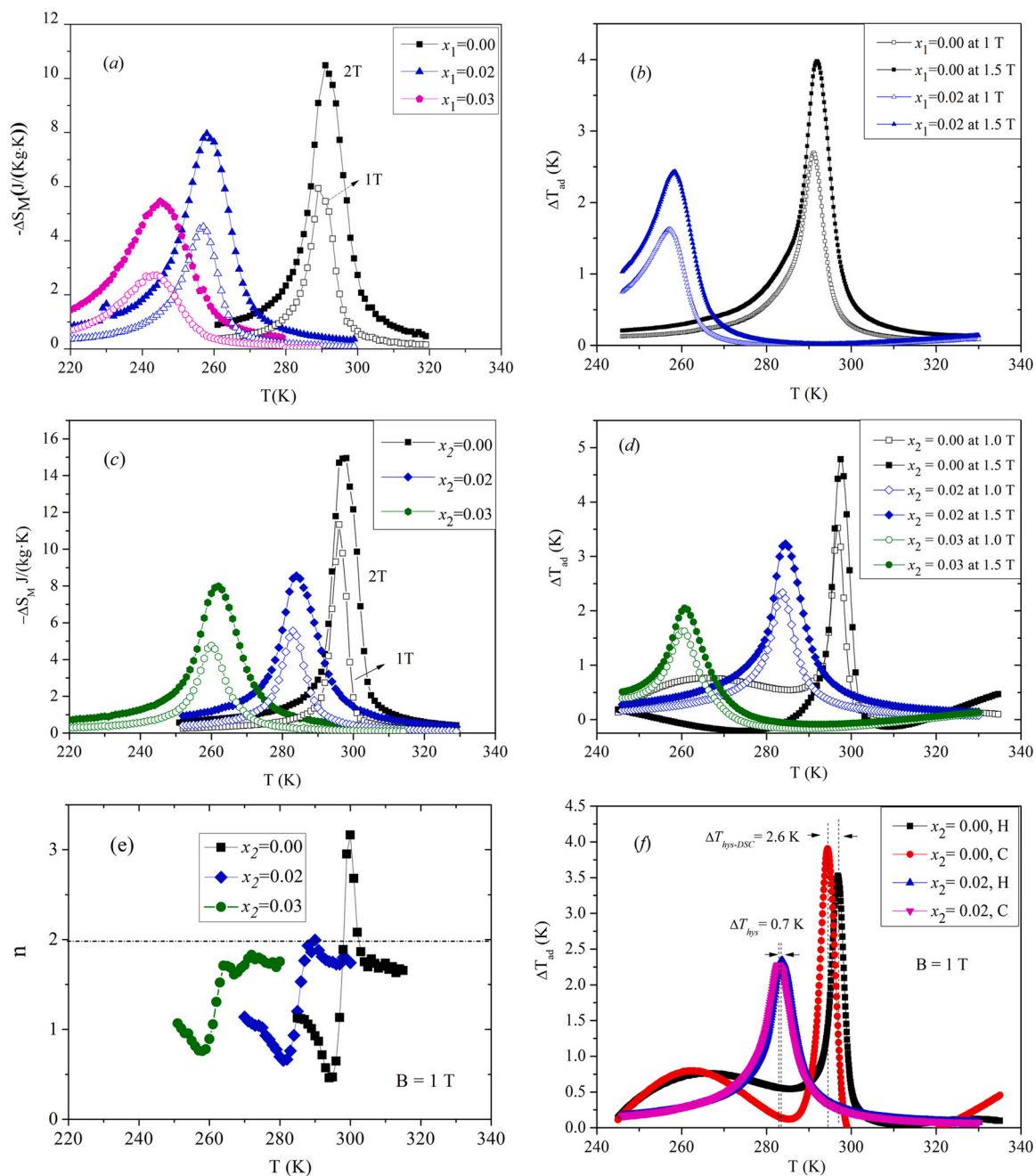
$$\Delta T_{ad} = - \left[ \frac{T}{C(T)_{H_F}} S_M(T)_{H_F} - \frac{T}{C(T)_{H_i}} S_M(T)_{H_i} \right] \quad (1)$$

Fig. 4(b) illustrates the temperature dependence of the in-field DSC values of  $\Delta T_{ad}$  for series A ( $x_1 = 0.00$  and 0.02) while Fig. 4(d) illustrates the temperature dependence of  $\Delta T_{ad}$  for series B ( $x_2 = 0.00, 0.02$  and 0.03). In the series A, when  $x_1$  increases from 0.00 to 0.02 at %, the value of  $\Delta T_{ad}$  decreases from 2.7 to 1.6 K under an applied magnetic field of 1 T. This value of  $\Delta T_{ad}$  for the benchmark Mn-Fe-P-Si-B [11] was 2.5 K. In the series B, when  $x_2$  increases from 0.00 to 0.02, its value of  $\Delta T_{ad}$  is decreased from 3.5 to 2.3 K at 1 T. At 1.5 T, the value of  $\Delta T_{ad}$  for the sample with  $x_2 = 0.02$  is 3.2 K. Thus,  $\Delta T_{ad}$  for the sample of  $x_2 = 0.02$  is sufficiently high to be utilized in a device, since cooling (and heating) may be effective when one achieves  $\Delta T_{ad}$  above 2.0 K [19].

The question is why the low hysteresis and high entropy change occurs in the sample with  $x_2 = 0.02$ . A quantitative criterion to determine the order of magnetic phase transitions through magnetocaloric curves has been proposed recently, i.e. the field exponent  $n = d \ln(|\Delta S_M|) / d \ln(H)$  for the relative field dependence of magnetic entropy change shows a maximum of  $n > 2$  near the transition



**Fig. 3.** Heating DSC curves of series A and B are measured under zero field.



**Fig. 4.** (a) and (c) Temperature dependence of  $|\Delta S_M|$  under a field change of 0–1.0 T (open symbols) and 0–2.0 T (filled symbols) for series A and B, respectively; (b) and (d) Temperature dependence of  $\Delta T_{ad}$  under a field change of 0–1.0 T (open symbols) and 0–1.5 T (filled symbols) during heating for series A and B, respectively; (e) Temperature dependence of the field exponent  $n = d \ln(|\Delta S_M|) / d \ln(H)$  under a field change of 1.0 T for  $x_2 = 0$  and 0.02; (f) Temperature dependence of  $\Delta T_{ad}$  under a field change of 0–1.0 T during heating and cooling for series B.

temperatures for a FOMT, while the absence of such overshoot indicates a SOMT [39,40]. For the critical composition, the minimum value of  $n$  corresponds to 0.4 [40–42]. The temperature dependence of the field exponent  $n$  under a field change of 1.0 T for  $x_2 = 0$  and 0.02 is shown in Fig. 4(e). For  $x_2 = 0$  sample, a clear distinct overshoot of  $n > 2$  is observed near its transition temperature, which indicates it is a FOMT. Due to the presence of impurity (3:1) phase (observed from XRD results) in the samples, which itself exhibits  $T_C = 575$  °C, [43]  $n$  decreases to values below 2 after the overshoot (the typical  $n$  value for paramagnetic state is 2) due to the ferromagnetic state of the existing (3:1) phase. It corresponds to the composition nearest to critical point of the FOMT and SOMT for  $x_2 = 0.02$ , since it is in good agreement with the shape of  $n$  for critical point proposed in

reference [39]. However, the presence of impurities can affect the absolute value of  $n$ , decreasing below the expected value of 2 for impurity phases with larger Curie temperatures. The observed slight increase of  $n$  to values close to 2 followed by a decrease might be an indication of a combined effect of a weakly first order phase transition and the presence of such impurities, which agrees with microstructural observations and the presence of a small thermal hysteresis. Accordingly, a good combination of low hysteresis and high entropy change can be obtained in this compound. For  $x_2 = 0.03$ , the minimum of  $n$  near the transition is rather shallow and two peaks are shown when passing the transition temperature. This type of shape for the field exponent  $n$  indicates that it has phase separation of  $Fe_2P$ -type structure, which agrees with the two peaks of

zero-field DSC in the Fig. 3. As shown in the Fig. S2, the phase separation of the Fe<sub>2</sub>P-type phase is observed for  $x_2 = 0.03$ . There are two peaks in the DSC curves, which show an interval of transition temperature about 1 K. The zero-hysteresis is shown in the M-T curve of  $x_2 = 0.03$  but is not shown for its in-field DSC data. The reason for the different value of hysteresis for M-T and DSC measurement is not clear yet. Recently, the origins for zero-hysteresis in the materials with IEM are investigated from the aspect of nanoscale phase separation [6,44,45] and the secondary phases [46]. Thus, we propose an explanation based on the in-field DSC data. Note that the  $T_{peak}$  of heating DSC curve for phase 1 and cooling DSC curve for phase 2 are overlapped. When the paramagnetic phase is transferred to ferromagnetic phase, the phase 1 with higher  $T_c$  will become ferromagnetic first and then the stray field [46] from this ferromagnetic phase can assist the phase transition of paramagnetic phase 2. As a result, the phase separation helps to achieve zero-hysteresis in M-T curves. Nevertheless, the phase separation may not be a good strategy for get a GMCE since it will also scarify the  $|\Delta S_M|$  and  $\Delta T_{ad}$ , see Fig. 4 (c) and (d) and shows hysteresis in the DSC measurement.

In addition,  $\Delta T_{hys-DSC}$  is determined by the difference of the heating and cooling process for the in-field DSC under a field change of 1 T measured at a rate of 3 Kmin<sup>-1</sup>, as shown in Fig. 4(f). It differs from the  $\Delta T_{hys}$  calculated by the M-T curves at a rate of 2 Kmin<sup>-1</sup>. Due to the thermal-lag effect, a higher measuring rate leads to a higher  $\Delta T_{hys}$ , as demonstrated for the Eu<sub>2</sub>In compound [47]. In series B,  $\Delta T_{hys-DSC}$  decreases from 2.4 to 0.7 K when  $x_2$  increases from 0.00 to 0.02,

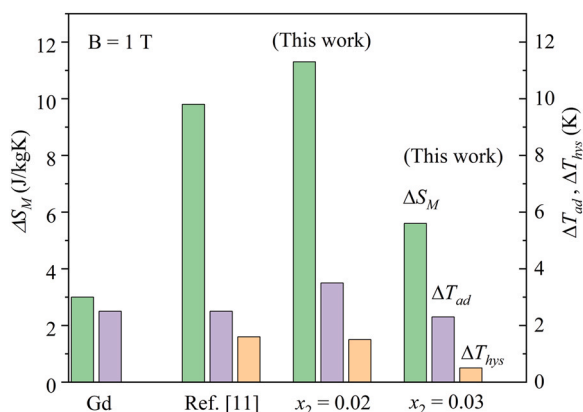


Fig. 5. Figure of merit of  $|\Delta S_M|$ ,  $\Delta T_{ad}$  and  $\Delta T_{hys}$  for Gd [18], MnFe<sub>0.95</sub>P<sub>0.563</sub>Si<sub>0.36</sub>B<sub>0.077</sub> [11], MnFe<sub>0.95</sub>P<sub>0.563</sub>Si<sub>0.36</sub>B<sub>0.077</sub> and Mn<sub>0.98</sub>V<sub>0.02</sub>Fe<sub>0.95</sub>P<sub>0.563</sub>Si<sub>0.36</sub>B<sub>0.077</sub> in series B.

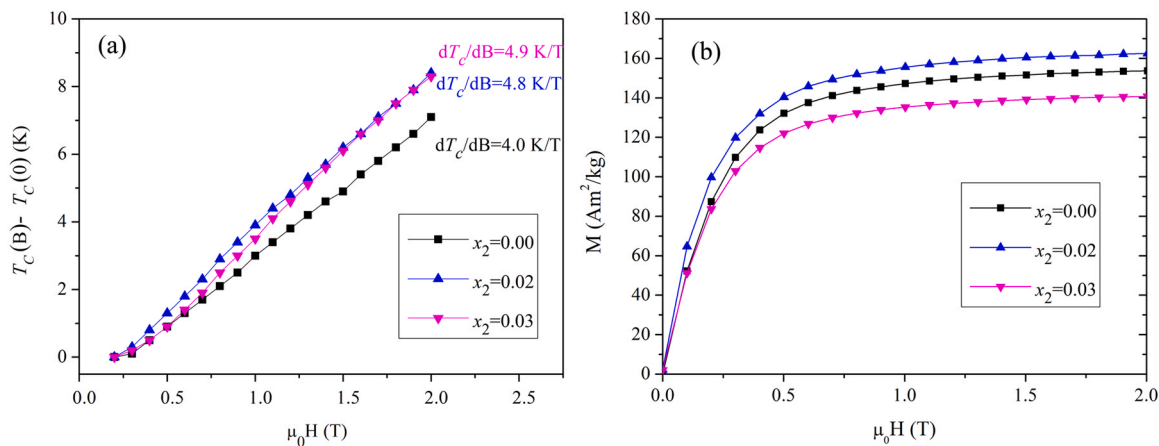


Fig. 6. Field dependence of  $T_c$  and  $dT_c/dB$  (insets); (b) The magnetization as a function of the V content for series B measured at a temperature of 5 K.

see Fig. 4(f). The value of  $\Delta T_{ad}$  for Mn<sub>0.98</sub>V<sub>0.02</sub>Fe<sub>0.95</sub>P<sub>0.563</sub>Si<sub>0.36</sub>B<sub>0.077</sub> ( $\Delta T_{ad} = 2.3$  K) in series B is competitive to that of MnFe<sub>0.95</sub>P<sub>0.563</sub>Si<sub>0.36</sub>B<sub>0.077</sub> compound ( $\Delta T_{ad} = 2.5$  K) [11], but its value for  $\Delta T_{hys-DSC}$  is reduced by 85%. Thus, it is promising to achieve a large value of  $\Delta T_{ad}$  and an ultra-low  $\Delta T_{hys-DSC}$ , which can significantly improve the heat exchange efficiency of the magnetic cooling (and heating) system.

The summary of maximum values for  $|\Delta S_M|$ ,  $\Delta T_{ad}$  and  $\Delta T_{hys}$  at a magnetic field of 1.0 T for Gd, MnFe<sub>0.95</sub>P<sub>0.563</sub>Si<sub>0.36</sub>B<sub>0.077</sub>, MnFe<sub>0.95</sub>P<sub>0.563</sub>Si<sub>0.36</sub>B<sub>0.077</sub> (this work) and Mn<sub>0.98</sub>V<sub>0.02</sub>Fe<sub>0.95</sub>P<sub>0.563</sub>Si<sub>0.36</sub>B<sub>0.077</sub> (this work) is shown in Fig. 5, demonstrating the merit of the compounds in this work. The value of  $\Delta T_{ad}$  for  $x_2 = 0.02$  is 2.3 K, comparable to 2.8 K of polycrystalline Gd, but its  $|\Delta S_M|$  is almost twice larger than Gd [18]. Accordingly, the coefficient of refrigerant performance ( $RCP = \frac{\Delta S^* \Delta T_{ad}}{\int_0^B M(T_c, B) dB}$ ) of the sample with  $x_2 = 0.02$  is calculated to be 0.47, larger than Gd ( $RCP = 0.17$ ) and Fe<sub>0.49</sub>Rh<sub>0.51</sub> ( $RCP = 0.22$ ) [48]. Thus, the MnFe<sub>0.95</sub>P<sub>0.563</sub>Si<sub>0.36</sub>B<sub>0.077</sub> ( $x_2 = 0.00$ ) and Mn<sub>0.98</sub>V<sub>0.02</sub>Fe<sub>0.95</sub>P<sub>0.563</sub>Si<sub>0.36</sub>B<sub>0.077</sub> ( $x_2 = 0.02$ ) are clearly good candidates for magnetic heat pumps operating near room temperature.

To investigate the reason why V decreases the hysteresis without losing the GMCE, the effect of V on  $dT_c/dB$  is investigated. The magnetic field dependence of  $T_c$  and  $dT_c/dB$  for series B are shown in Fig. 6(a). The value of  $dT_c/dB$  increases from 4.0 to 4.8 K/T when the V content is changed from  $x_2 = 0.00$  to  $x_2 = 0.02$ . This is comparable to the  $dT_c/dB$  value for (Mn,Fe)<sub>2</sub>(P,As) compounds, where  $dT_c/dB$  was found to be 5.2 K/T [49]. The Clausius–Clapeyron relation for a FOMT corresponds to  $dT_c/dB = -T_c \Delta M/L$ , where B is the applied magnetic field and  $\Delta M$  is the jump in magnetization. Thus, the increase in  $dT_c/dB$  is mainly caused by the decrease in latent heat (see Table 1), since the values of  $T_c$  and  $\Delta M$  are reduced (see Fig. 2). The increase in  $dT_c/dB$  suggests that the driving force for the FOMT is enhanced. As a result, a strong field response is found for low fields, as shown in the magnetization curve of  $x_2 = 0.02$  in the Fig. 6(b).

From the saturation magnetization displayed in Fig. 6(b), the value of the magnetic moment per formula unit ( $\mu_{f.u.}$ ) for series B was calculated, as mentioned in reference [50]. For series B, the value of  $\mu_{f.u.}$  increases from 3.75 to 3.96  $\mu_B/f.u.$  when  $x_2$  increases from 0.00 to 0.02. A larger value of  $\mu_{f.u.}$  suggests a larger contribution to  $|\Delta S_M|$ . This agrees with the strengthening effect of V substitution on the magnetic moment formation in the Mn-Fe-P-Si-V compounds [31]. The higher values for  $dT_c/dB$  and  $\mu_{f.u.}$  illustrate why an ultra-low thermal hysteresis and a GMCE can be achieved simultaneously in the compounds with 0.02 at% V substitution for Mn in series B. Thus, introducing V as a new substitutional element is found to be capable of increasing both  $dT_c/dB$  and  $\mu_{f.u.}$  and thereby decreasing the hysteresis without losing the GMCE.

## 4. Conclusions

The ultra-low hysteresis and GMCE of optimal  $Mn_{1-x_2}V_{x_2}Fe_{0.95}P_{0.563}Si_{0.36}B_{0.077}$  compounds pave a pathway to highly efficient magnetic heat pump applications. When  $x_2$  increases from 0.00 to 0.02, the latent heat can be reduced by 55% from 6.2 to 3.3 J/g and thus  $\Delta T_{hys}$  decreases by 67% from 1.5 to 0.5 K, the  $|\Delta S_M|$  is decreased from 11.3 to 5.6 J/kgK and the  $\Delta T_{ad}$  is decreased from 3.5 to 2.3 K at 1 T. The optimal composition of  $x_2 = 0.02$  is proven to be close to the critical point between the FOMT and the SOMT. The combination of increase in  $dT_C/dB$  and increase in magnetic moment  $\mu_{f.u.}$  of  $Mn_{0.98}V_{0.02}Fe_{0.95}P_{0.563}Si_{0.36}B_{0.077}$  compound can provide a sufficiently large GMCE and an ultra-low value (0.5 K) of  $\Delta T_{hys}$ . Thus, the coefficient of refrigerant performance (CRP) of this compound is two times larger than Gd. The current  $Mn_{1-x}V_xFe(P,Si,B)$  compounds provide an excellent candidate for designing materials towards magnetic heat pump applications using low-cost permanent magnets near-room temperature. The mechanical properties and thermal conductivity of the  $Mn_{1-x_2}V_{x_2}Fe_{0.95}P_{0.563}Si_{0.36}B_{0.077}$  compounds will need to be investigated further to demonstrate the full potential for future utilization in practical devices.

## CRedit authorship contribution statement

**Ekkas Brück:** Conceptualization, Methodology, Reviewing. **Jiawei Lei:** Data curation, Writing – original draft. **Xinmin You:** Data curation, Writing. **Jiayan Law:** Data curation, Writing. **Victorino Franco:** Data curation, Writing. **Bowei Huang:** Data curation, Writing. **Dimitrios Bessas:** Data curation, Writing. **Michael Maschek:** Data curation, Writing. **Dechang Zeng:** Supervision, Methodology, Reviewing. **Niels van Dijk:** Visualization, Investigation, Writing – review & editing.

## Data availability

Data will be made available on request.

## Declaration of Competing Interest

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests: Jiawei Lai, Victorino Franco reports financial support was provided by Guangzhou ethics project. Michael Maschek reports financial support was provided by EIT Climate KIC. Ekkas Brück has patent #P1600096NL00 pending to Technische Universiteit Delft. Jiawei Lai has patent #P1600096NL00 pending to Technische Universiteit Delft.

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## Appendix A. Supporting information

Supplementary data associated with this article can be found in the online version at doi:10.1016/j.jallcom.2022.167336.

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