Robust Bain distortion in the premartensite phase of a platinum-substituted Ni$_2$MnGa magnetic shape memory alloy

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The premartensite phase of shape memory and magnetic shape memory alloys (MSMAs) is believed to be a precursor state of the martensite phase with preserved austenite phase symmetry. The thermodynamic stability of the premartensite phase and its relation to the martensitic phase is still an unresolved issue, even though it is critical to the understanding of the functional properties of MSMAs. We present here unambiguous evidence for macroscopic symmetry breaking leading to robust Bain distortion in the premartensite phase of 10% Pt-substituted Ni$_2$MnGa. We show that the robust Bain-distorted premartensite (T2) phase results from another premartensite (T1) phase with preserved cubic-like symmetry through an isostructural phase transition. The T2 phase finally transforms to the martensite phase with additional Bain distortion on further cooling. Our results demonstrate that the premartensite phase should not be considered as a precursor state with the preserved symmetry of the cubic austenite phase.
The appearance of a precursor state, widely known as the premartensite (PM) phase, is an interesting feature of compounds/alloys exhibiting martensitic phase transitions. This has been extensively studied in conventional shape memory alloys (SMAs), where shape recovery can be realized under the influence of not only the temperature and stress but also the magnetic field. Among the SMAs, stoichiometric Ni$_2$MnGa and off-stoichiometric Heusler alloys in the alloy systems Ni–Mn–X (Ga, In, Sn, Sb) have received considerable attention. The stoichiometric Ni$_2$MnGa system is by far the most studied due to its large (10%) magnetic field-induced strain (MFIS) with potential for applications in novel multifunctional sensor and actuator devices. The recent discovery of skyrmions in Ni$_2$MnGa has opened newer potential applications for this material in the field of spintronics as well. Ni$_2$MnGa is a multiferroic SMA with two ferroic order parameters, namely spontaneous magnetization and spontaneous strain, which appear below the ferromagnetic and ferroelastic (martensite) phase transition temperatures $T_C \approx 370$ K and $T_M \approx 210$ K, respectively. The martensitic transition in Ni$_2$MnGa is preceded by a PM phase transition around $T = 260$ K. Both the martensite and PM phases exhibit very strong coupling between the two ferroic order parameters (i.e., magnetoelastic coupling) which is responsible for the large MFIS. The large MFIS is also closely linked with the existence of a long period modulated structure of the martensite phase in the Ni$_2$MnGa alloy. Since the modulated phase of Ni$_2$MnGa appears via a modulated PM phase and not directly from the austenite phase after its Bain distortion, understanding the characteristics of the PM phase and its effect on the martensite phase transformation has been a topic of intense research in recent years. The austenite-PM and the PM-martensite transitions are first order in nature, with characteristic thermal hysteresis and phase coexistence regions, due to the strong magnetoelastic coupling between the magnetization and strain caused by a soft $TA_2$ phonon mode at a wave vector $q \sim 1/331, 33, 39–41$. The nature of the modulation in the PM and martensite phases has been shown to be 3 M and 7M-like but incommensurate. The incommensurately modulated PM and the martensite phases coexist over a wide temperature regime across $TSM$, suggesting that the martensite phase in Ni$_2$MnGa originates from the PM phase through a first-order phase transition. While the martensite phase of Ni$_2$MnGa exhibits macroscopic symmetry breaking, the PM phase does not exhibit any evidence of symmetry breaking transition.

In the phenomenological theories of martensite phase transition, the transformation occurs through a lattice deformation shear, with atomic shuffles if necessary, leading to Bain distortion of the austenite unit cell while maintaining an invariant habit plane (contact plane) between the parent and the martensite phases achieved through a lattice invariant shear produced by twinning or stacking faulting. Obviously, it would have been kinetically more favorable if the austenite phase had undergone a mild Bain distortion in the PM phase that could have provided a further pathway to the martensite phase transformation with higher Bain distortion while maintaining the integrity of the habit plane all along. In the absence of any macroscopic Bain distortion, the PM phase of Ni$_2$MnGa does not seem to have a role in facilitating the appearance of Bain distortion in the martensite phase with an invariant habit plane. This raises a fundamental question as to why at all the PM phase appears in Ni$_2$MnGa. Interestingly, first principles calculations reveal a local minimum that corresponds to an intermediate value of the Bain distortion in the PM phase. However, this has not been verified experimentally till now.

We present here experimental evidence for macroscopic symmetry breaking in the PM phase of Ni$_2$MnGa as a result of 10% Pt substitution leading to a robust Bain distortion. From temperature-dependent high resolution synchrotron X-ray powder diffraction (SXRPD) data analysis, we show that the Bain distortion grows in steps. First, a nearly 3 M modulated PM phase (PM(T1)) with an average cubic-like feature (i.e., negligible Bain distortion) of the elementary L2$_1$ unit cell results from the austenite phase. This phase then undergoes an isorotational phase transition to another 3M-like PM phase (PM(T2)) with robust Bain distortion at lower temperatures. Our results reveal that the two PM phases are stable thermodynamic phases and not precursor phases with preserved austenite symmetry, as has been presumed in the literature all along. Further, we also show that the martensite phase originates from the larger Bain-distorted PM phase suggesting that the Bain distortion appears in steps to facilitate the emergence of an invariant habit plane. These observations provide us an opportunity to critically examine the two existing mechanisms of the origin of modulation in these systems.

**Results**

**Temperature-dependent ac-susceptibility.** The temperature ($T$) dependence of the ac-susceptibility ($\chi(\omega)$) of the Ni$_{2-x}$Pt$_x$Ni$_2$MnGa powder sample shown in Fig. 1 reveals a sharp change at paramagnetic-ferromagnetic transition temperature $T_C \approx 370$ K. On further cooling, the ac-$\chi$ decreases with a change of slope in the $\chi$–$T$ plot around $T \approx 260$ K. Interestingly, there is another change of slope in ac-$\chi$ around $T = 235$ K preceded by a large (~40%) drop in ac-$\chi$. In general, a large drop in ac-$\chi$ (or dc magnetization in low field) in the Ni-Mn-Ga based SMAs is
related with the large magnetocrystalline anisotropy of the martensite phase\textsuperscript{27, 46} but Fig. 1 suggests that nearly 40% of the total drop in ac-χ has already occurred at 235 K in the PM phase as a result of 10% Pt substitution. It is important to mention here that while in Ni\textsubscript{2}MnGa a very small (nearly 0.5%) drop in ac-χ at the PM phase during cooling from the austenite phase is observed\textsuperscript{27} for Ni\textsubscript{1.9}Pt\textsubscript{0.1}MnGa this drop is very large, which indicates a larger magnetocrystalline anisotropy of the PM phase in Ni\textsubscript{1.9}Pt\textsubscript{0.1}MnGa.

**High resolution synchrotron X-ray powder diffraction.** Our high resolution SXRPD patterns show clear correlation between the change of slope in the ac-χ-T plot and structural changes corresponding to the PM and martensite phases discussed above. We show in Fig. 2 the variation of SXRPD patterns in selected 2θ ranges with temperature in the 290–110 K range which captures the entire sequence of phase transitions. The SXRPD pattern at 290 K corresponds to the cubic austenite phase as it does not show any splitting of the XRD peaks. The Rietveld refinement using the 290 K data for the cubic structure (Fm-3m space group) shows excellent fit between the observed and calculated peak profiles which further confirms the cubic structure at this temperature (Fig. 3a). The refined cell parameter a = 5.84753(6) Å obtained from the Rietveld refinement is slightly higher than that of the stoichiometric Ni\textsubscript{2}MnGa (a = 5.82445(1) Å), which might be due to the larger atomic size of the Pt (1.77 Å) atoms compared to the Ni (1.37 Å) atoms. This is in good agreement with an earlier study\textsuperscript{46}. On cooling the sample below T = 260 K, several smaller intensity peaks appear. Two such peaks are marked in Fig. 2b and the inset of Fig. 2b. The low intensity peaks are the satellites that appear due to the modulated nature of the PM phase in Ni\textsubscript{1.9}MnGa\textsuperscript{27}. The intensity of these satellites increases without much effect on the austenite cubic peaks on further cooling up to T = 240 K. The situation is similar to that in Ni\textsubscript{2}MnGa, where cooling below the PM phase transition temperature (T\textsubscript{PM} = 261 K) leads to the appearance of low intensity peaks. On the other hand, the austenite peaks remain nearly unaffected and retain their cubic nature\textsuperscript{27}. The SXRPD of Ni\textsubscript{1.9}Pt\textsubscript{0.1}MnGa in the temperature range 260–240 K thus looks similar to the modulated structure of the PM phase of Ni\textsubscript{1.9}MnGa, which is incommensurate with 3M-like modulation\textsuperscript{25–28, 30, 30, 51}. Therefore, we carried out Rietveld analysis of the 240 K SXRPD pattern of Ni\textsubscript{1.9}Pt\textsubscript{0.1}MnGa using the (3 + 1) D superspace group approach taking into account both the main and the satellite reflections. We use the modulated structure of the PM phase similar to that of Ni\textsubscript{2}MnGa with the superspace group Immm (00y) y00. A fit between the observed and calculated peak profiles can be seen in Fig. 3b. The refined lattice parameters at 240 K are found to be: a = 4.1337(2) Å, b = 5.8416 (3) Å, and c = 4.1325 (1) Å with a modulation wave vector \( \mathbf{q} \) of \( q = 0.325 \mathbf{c}^* = (1/3 - \delta) \mathbf{c}^* \), where \( \delta = 0.0083 \) is the degree of incommensuration of modulation at 240 K. The magnitude of \( \mathbf{q} \) confirms the incommensurate nature of the modulation. The closest rational approximant to the observed value is (1/3)\( \mathbf{c}^* \) which suggests 3M-like modulation in the PM phase. Thus the modulated structure of the PM phase of Ni\textsubscript{1.9}Pt\textsubscript{0.1}MnGa stable in the temperature range 260–240 K, is identical to the PM phase of Ni\textsubscript{1.9}MnGa. The refined lattice parameters and modulation vector are also close to the PM phase of Ni\textsubscript{2}MnGa\textsuperscript{25}.

At ~235 K, which is close to the slope change in the ac-χ data (marked with the tick in Fig. 1), an interesting feature is observed in the diffraction pattern. At this temperature, the splitting of the Bragg peaks of the cubic austenite structure begins to show (Fig. 2d, e). However, no additional/new satellite reflections...
We investigated the structure of the martensite phase at 110 K by Rietveld refinement in the \((3 + 1)\) D superspace similar to that in Ni_2MnGa. The main peaks originating from the splitting of the main austenite peaks could be indexed using an orthorhombic unit cell and space group \(Immm\) as in case of Ni_2MnGa. The unit cell parameters obtained after Le Bail refinement were found to be \(a = 4.2390\ (1)\ \text{Å}, \ b = 5.5682\ (1)\ \text{Å},\) and \(c = 4.2074\ (1)\ \text{Å}.\) The Rietveld refinement was then carried out using the superspace group \(Immm\) \((00\gamma)\) s00 by taking into account the complete diffraction pattern including both the main and satellite reflections. The good fit between the observed and calculated profiles (Fig. 3c) confirms that the refinement has converged successfully. The refined value of the incommensurate modulation vector was found to be \(q = 0.4290\ (2)\ c^* = (3/7 + \delta)\ c^*,\) where \(\delta = 0.00112\) is the degree of incommensuration. Thus the structure at 110 K (which is a representative of the martensite phase stable below 220 K) shows incommensurate modulation similar to the martensite phase of Ni_2MnGa\(^{27, 28, 42, 52}\), which is of nearly 7M type.

The structures of the cubic austenite, PM in the temperature range 260–240 K and martensite at 110 K, which is representative of the structure below 220 K of Ni$_{1+\delta}$Pt$_{0.1}$MnGa, are similar to Ni$_2$MnGa. However, the structure of the PM phase of Ni$_{1+\delta}$Pt$_{0.1}$MnGa in the temperature range 235–195 K, over which the splitting of the cubic austenite peaks is observed without the presence of the martensite peaks, is quite unusual and we now proceed to analyze this newly observed phase. The splitting of the austenite peaks in the PM phase indicates the loss of pseudocubic symmetry of the basic structure. This is unexpected for the PM phase as it is always considered to be a micromodulated phase with preserved cubic symmetry of the austenite peaks. We consider the 225 K data as a representative of this new structure and present the result of fits between the calculated and observed intensities by Rietveld refinement. As in the case of the Rietveld refinement of the PM and martensite phases stable in the temperature ranges 245 K < \(T < 260\) K and \(T < 225\) K, respectively we first considered only the main reflections (not satellites) and carried out Le Bail refinement using the cubic \(Fm-3m\) space group. From Fig. 4a, it can be clearly seen that this model misses out the new peaks, which result due to the splitting of the cubic austenite peaks (see the insets of Fig. 4a). This confirms that the cubic symmetry is now broken. We therefore considered the tetragonal unit cell with space group \(I4/mmm\), which could capture the splitting but still the observed and calculated peak profiles are not matched properly. Finally, we took orthorhombic distortion into account in the space group \(I4/mmm\), which was also used for the martensite phase. The excellent fit between observed and calculated peak profiles (Fig. 4c and inset) shows that at 225 K the basic structure is orthorhombic with refined lattice parameters as: \(a = 4.1356\ (2)\ \text{Å}, \ b = 5.8276\ (2)\ \text{Å},\) and \(c = 4.1371\ (1)\ \text{Å}.\) There is a clear evidence for the pseudo-tetragonal Bain distortion of the PM phase with \(b/\sqrt{2a} \sim 0.9965.\) Having established the Bain-distorted orthorhombic nature of the PM phase at 225 K, we then carried out Rietveld refinement considering the complete diffraction pattern including satellite reflections using superspace group \(I4/mmm\) \((00\gamma)\) s00. A good fit between the observed and calculated peak profiles was obtained after the Rietveld refinements (Fig. 4d and inset). The refined modulation wave vector \(q = 0.3393\ c^*\) shows incommensurate nature of the modulation with \(q = (1/3 + \delta)\ c^*,\) where \(\delta = 0.0059\) is the degree of incommensuration. Our results using high resolution SXRD data thus provide evidence for a new incommensurately modulated 3M-like PM phase with broken cubic austenite symmetry. This is the example of macroscopic symmetry breaking and Bain distortion in the PM phase of any Heusler MSMA.
After observing a different type of the PM phase with robust Bain distortion, we now proceed to present evidence for the coexistence of the incommensurate 3M-like Bain-distorted PM and incommensurate 7M-like martensite phases below 220 K. For this, we show in Fig. 5 the results of Rietveld refinement using the SXRPD pattern at 210 K. It can be observed from the inset of Fig. 5a that consideration of only the 3M-like incommensurate PM phase cannot account for some of the Bragg reflections (marked with M in the inset of Fig. 5a). We also verified that single phase 7M-like incommensurate martensite phase structure cannot account for this diffraction pattern. However, a refinement based on coexistence of both 3M-like Bain-distorted PM as well as 7M-like martensite phases, both with incommensurate modulations, coexist at 210 K. Using superspace Rietveld refinement, the coexistence of 3M-like and 7M-like incommensurate structures was confirmed in the entire temperature range from 210 to 195 K. As phase coexistence is a typical characteristic of a first-order phase transition, our result show that the Bain-distorted PM phase to martensite phase transition is a first-order transition. Hence, the martensite phase originates from the PM phase in Ni$_{1.9}$Pt$_{0.1}$MnGa similar to that in Ni$_2$MnGa$^{27}$. However, there is a major difference in the case of Ni$_{1.9}$Pt$_{0.1}$MnGa since the PM phase from which the martensite phase results shows robust Bain distortion in marked contrast to Ni$_2$MnGa where the martensite phase results from a PM phase that preserves the cubic symmetry of the main unit cell.

Figure 6a shows the variation of the lattice parameters and unit cell volume for the equivalent cubic, PM ($a_{pm} \approx (1/\sqrt{2}) \ a_c$, $b_{pm} \approx a_c$, $c_{pm} \approx (1/\sqrt{2}) \ a_c$) and martensite ($a_{m} \approx (1/\sqrt{2}) \ a_c$, $b_{m} \approx a_c$, $c_{m} \approx (1/\sqrt{2}) \ a_c$) with temperature. It is evident from Fig. 6a that unit cell volume for all the phases decreases with decreasing temperature. However, the volume of the cubic austenite phase changes rather smoothly across the austenite to PM (T1) phase transition (260–245 K), similar to that in Ni$_2$MnGa, as expected for a weak first-order phase transition. However, on lowering the temperature further, a discontinuous change in the volume is observed at T $\approx$ 235 K with a concomitant splitting of the cubic austenite peaks (Figs. 6a and 2e) and change of slope in the ac-$\chi$ vs. T plot (Fig. 1). This discontinuous change in volume occurs at the cubic martensite to the Bain-distorted DM phase transition and confirms the first-order nature of this phase transition. Furthermore, a clear discontinuous change in volume occurs during Bain-distorted PM (T2) to martensite phase transition is also observed confirming the first-order character of
this transition as well. The $b/a$ ratio shown in the inset of Fig. 6a also exhibits discontinuous change at temperatures corresponding to the different phases. We believe that the formation of the PM (T1) phase in the cubic austenite matrix, followed by robust Bain-distorted PM (T2) phase in the PM (T1) matrix and finally the fully Bain-distorted martensite phase in the PM (T2) phase is kinetically more favorable for maintaining an invariant habit plane all through the transition due to the gradual evolution of the Bain distortion.

The displacement of atoms obtained from the Rietveld refinements (Supplementary Discussion and Supplementary Table 1) for PM(T1) at 240 K and PM (T2) at 225 K phases are compared in Fig. 6c–e, which clearly indicates a change in the atomic displacement between two phases. Since, both the PM phases belong to the same superspace group $I\overline{4}mm$ (000) s00 with nearly 3M-like incommensurate modulation, this is an isostructural phase transition where the atomic positions shift without changing the Wyckoff site symmetries and the over all space group. The 3D approximant structures of PM (T1) and PM (T2) phases are compared in Supplementary Table 2. The discontinuous change in the unit cell parameters at the isostructural phase transition temperature clearly reveals the presence of strong spin lattice (magnetoelastic) coupling. Isostructural phase transitions are rather rare and have been reported for example in the past during electronic transitions in chalcogenides under high pressure, in multiferroics across the magnetic phase transition leading to excess spontaneous polarization due to magnetoelectric coupling, between two ferroelectric phases and between two antiferrodistorted structures in CaTiO$_3$. Our results provide evidence for an isostructural phase transition from a nearly cubic-like PM phase to a robust Bain-distorted PM phase in a magnetic SMA.

An incommensurate phase is often perceived to undergo a transition to the lock-in commensurate phase at low temperatures in the ground state. However, our recent high resolution SXRPD study on the stochiometric Ni$_2$MnGa composition reveals that modulation wave vector ($\mathbf{q}$) as a function of temperature using SXRPD data down to 110 K. In addition, we also carried Le Bail refinement using high resolution neutron powder diffraction data at 10 K to verify whether the incommensurate phase would lock-into a commensurate phase in the ground state. The temperature variation of the modulation wave vector $\mathbf{q}$ is shown in Fig. 6b during cooling. A jump $q$ (magnitude of $\mathbf{q}$) is clearly observed at the 3M modulated PM (T1) to PM (T2) isostructural phase transition while both phases continue to...
that the volume expansion due to Pt substitution causes the robust Bain distortion in Ni$_{1.83}$Pt$_{0.17}$MnGa.

To further check this, we calculated energies as function of $c/a$ ratios for Ni$_2$MnGa at the two different volumes. Our calculations reveal again an energy plateau with reduced value for Ni$_2$MnGa at its equilibrium volume without an energetic preference for $c/a \sim 0.993$ (blue circles in Fig. 8b), but does not exclude the possibility to observe the Bain-distorted phase in unsubstituted composition. However, at the larger equilibrium volume of Ni$_{1.83}$Pt$_{0.17}$MnGa, the Bain-distorted PM state has lower energy at $c/a \sim 0.993$, as can be clearly seen from Fig. 8b. Interestingly, we find that the reduced magnetization at finite temperature captured here through fixed-spin moment calculations stabilizes the cubic PM phase (Fig. 8b). Hence, while the Pt substitution at the Ni site modifies the energy plateau found for Ni$_2$MnGa through the enhanced volume and stabilizes the robust Bain-distorted PM phase at lower temperatures, the finite temperature decrease in magnetization will stabilize the cubic PM phase at higher temperatures. This explains the experimentally observed sequence of structural transitions in the PM phase.

In Fig. 8c, we present calculated magnetic moments for Ni$_2$MnGa at the two different volumes. It can be clearly seen that the magnetic moments slightly (by ~1%) increase at the higher volume of Ni$_{1.83}$Pt$_{0.17}$MnGa (orange squares) as compared to those obtained at the equilibrium volume of Ni$_2$MnGa (blue circles). The more important observation is a discontinuous jump in magnetic moment for $c/a \sim 0.993$ at higher volume. This discontinuous jump is linked with the sudden change of the energy state at the same $c/a$ ratio (Fig. 8b).

Discussion

In this work, we have presented evidence for robust Bain distortion in the PM phase of Ni$_2$MnGa as a result of 10% Pt substitution using high resolution SXRPD. This finding is significant for the field of SMA in general and SMAs in particular. Our results show that the austenite to martensite phase transition in Ni$_{1.83}$Pt$_{0.17}$MnGa involves three intermediate steps. The first transition is from the austenite to 3M-like incommensurate modulated PM phase (PM (T1)), which preserves the cubic symmetry of the austenite peaks similar to that in the stoichiometric Ni$_2$MnGa. The second transition is an isostructural transition from 3M-like incommensurate modulated PM phase (PM (T1)) with negligible Bain distortion to another 3M-like incommensurate modulated PM phase (PM (T2)) with robust Bain distortion without any change in the superspace group symmetry. Then this new PM (T2) phase finally transforms to the 7M-like incommensurate modulated martensite (M) phase. Our results thus suggest that Pt substitution leads to gradual increase in Bain distortion which can facilitate the invariant habit plane requirement all through the three transitions until the martensite phase is formed. Our first principles calculations show that volume expansion due to Pt substitution provides the driving force for the robust Bain distortion of the PM (T2) phase in Pt doped Ni$_2$MnGa at lower temperatures while the reduced magnetization stabilizes the cubic-like PM (T1) phase at higher temperatures. Our findings provide an opportunity to critically reevaluate the applicability of the two main theories suggested for the origin of the modulation. The first one is related to the adaptive phase concept, which considers the ground state of the martensite phase to be L1$_0$ type non-modulated tetragonal structure resulting from the lattice deformation strain (Bain distortion) of the cubic austenite phase. In this model the modulated structure is a metastable phase formed due to kinetic reasons through nanotwinning of the Bain-distorted austenite structure. This model does not support the existence of the intermediate PM phase and treats the martensite

Fig. 7 Le Bail fits for the neutron diffraction patterns at 10 K. The experimental data, fitted curve and the residue are shown by circles (black), red continuous line, and green continuous line, respectively. The tick marks (blue) represent the Bragg peak positions

Theory. To understand the role of Pt in causing robust Bain distortion in the PM phase, we have performed first principles calculations of the stoichiometric Ni$_2$MnGa and Ni$_{1.83}$Pt$_{0.17}$MnGa (shown in Fig. 8). For Ni$_{1.83}$Pt$_{0.17}$MnGa (orange squares in Fig. 8a), our calculations reveal a pronounced energy minimum at $c/a \sim 0.993$, which indicates Bain distortion in the PM phase of this composition. This Bain distortion is caused by Pt, since a similar minimum is not present in Ni$_2$MnGa (blue circles in Fig. 8b). Since Pt is non-magnetic, it is unlikely to cause direct magnetoelastic coupling, which is otherwise typical for NiMn-based Heusler alloys. Instead, both our measurements and calculations indicate that the substitution of larger Pt atoms in place of smaller Ni atoms increases the optimum volume of the alloy. In order to clearly understand the role of volume, we have calculated the total energy of Ni$_{1.83}$Pt$_{0.17}$MnGa as function of $c/a$ ratio at the lower optimum volume of Ni$_2$MnGa (blue circles in Fig. 8a). It can be seen that lowering the volume yields a flat energy plateau with no clear minimum present at $c/a \sim 0.993$. This clearly shows
phase as a kinetically stabilized state to overcome the lattice deformation shear causing the Bain distortion. The other theory is based on a displaceable modulation concept, where the origin of modulation is related to a TA2 soft acoustic phonon mode of the austenite phase that gets coupled to the charge density wave in the PM phase with opening of a pseudogap at the Fermi surface. In this model, an incommensurate modulated martensite phase can result from the incommensurate modulated PM phase, both of which are stable thermodynamic phases in their respective range of temperatures. Our results clearly show that the incommensurate martensite phase with larger Bain distortion appears in steps from the Bain-distorted PM phase, which supports the soft phonon mode based model as the likely origin of modulation in Ni$_2$MnGa. The understanding of the origin of modulation in these alloys provides a pathway to design new MSMAs.

Before we close, we would like to briefly mention the likely implications of our results on the functional properties of Ni$_2$MnGa. In view of the fact that large MFIS observed in Ni$_2$MnGa is intimately linked with the existence of the modulated structure, the gradual evolution of modulation wave vector in going from PM (T1) to PM (T2) to martensite phase may influence the magnitude of MFIS. MSMAs also exhibit magnetocaloric effect (MCE) but the off-stoichiometric Ni–Mn–X (X = Ga, In, Sn, Sb) show larger MCE as compared to Ni$_2$MnGa. However, the off-stoichiometric alloys show larger hysteresis leading to irreversible MCE, which is a major drawback for any practical application. Since the gradual evolution of the Bain distortion can facilitate the emergence of an invariant habit plane, we believe that such alloys may exhibit better reversibility due to lower hysteresis.

Methods
Sample preparation and characterization. A polycrystalline sample of Ni$_{1.8}$Pt$_0.1$MnGa was prepared by the standard arc melting technique. The actual composition of the sample was checked by energy dispersive analysis of X-rays to be Ni$_{1.8}$Pt$_{0.1}$Mn$_{1.9}$Ga. The characteristic (magnetic and structural) transition temperatures were obtained by the ac-susceptibility measurements as a function of temperature using a SQUID-VSM magnetometer.

Synchrotron X-ray and neutron powder diffraction and analysis. High resolution synchrotron powder X-ray diffraction (SRPD) measurements were performed at the P02 beamline in Petra III, Hamburg, Germany using a wavelength of 0.20712 Å. The time of flight neutron diffraction data were obtained at the ISIS facility (UK) using the high resolution powder diffractometer HRPD at 10 K. For SRXD and neutron diffraction measurements, the powder samples obtained by crushing the as-cast ingots were annealed at 773 K under a high vacuum of $10^{-5}$ mbar for 10 h to remove the residual stresses introduced during grinding. The elastic wave vector of the magnetic structure was fixed and the data were collected using a fixed spin moment approach at 0 K with a $c/a$ ratio for the 3M premartensite phase of Ni$_{1.83}$Pt$_{0.17}$MnGa and Ni$_2$MnGa. Square symbols correspond to calculations performed at the equilibrium volume of Ni$_{1.83}$Pt$_{0.17}$MnGa, while circles denote calculations performed at the equilibrium volume of Ni$_2$MnGa. The energy computed with fixed-spin moment approach for Ni$_2$MnGa at the equilibrium volume of Ni$_{1.83}$Pt$_{0.17}$MnGa is shown by triangles. The variation of magnetic moment with $c/a$ ratio for Ni$_2$MnGa at its optimum volume (circles) and at the optimum volume of Ni$_{1.83}$Pt$_{0.17}$MnGa (squares). The red ellipse marks a discontinuous jump of the magnetization (yellow curve).

**Theoretical calculation.** For theoretical calculations, we have chosen a slightly different Pt concentration as compared to the experimental composition in order to minimize computational time. Nevertheless, we demonstrate that the physical mechanism responsible for the Bain distortion of the PM phase does not depend on the precise Pt concentration. Further, we have assumed a commensurate 3 M structure for the PM phase and have also neglected finite temperature entropic contributions. The spin polarized density functional theory (DFT) calculations have been performed for a 24-atom supercell using Vienna Ab-Initio Simulation Package. Relaxed geometries and atomic positions are determined using the projector-augmented wave method. The exchange-correlation functional is described by the generalized gradient approximation of Perdew–Burke–Ernzerhof. To avoid errors in calculated total energy because of artificial Pulay stress arising due to incomplete basis set, an energy cutoff of 700 eV is used for the plane wave basis and the independence of this choice (achieved convergence in the total energy of 0.01 meV/atom) has been checked. For the Brillouin zone sampling, a k-point grid of $8 \times 24 \times 18$ is used. Convergence criteria for the electronic structure and the largest residual forces of respectively $10^{-5}$ eV and $10^{-3}$ eV/Å are used. Such strict choice of cutoff parameters and convergence criteria resulted in DFT energies with error $<0.003$ meV/atom. Finite temperature magnetic effects have been modeled by constraining the total moment per cell by means of the fixed-spin moment approach.

**Data availability.** The data that support the findings of this study are available upon request from S.S.

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Author contributions
S.S. proposed the problem. C.F. and D.P. supervised the project. S.S. prepared the sample. P.D. performed compositional analysis. S.S. performed magnetic and synchrotron experiments. S.S., M.G.Z., and A.S.G. did neutron diffraction experiment. S.S. and D.P. analyzed the experimental data. B.D. performed theoretical calculations. S.W.D.S., S.C., and T.H. provided inputs for theory. S.S. and D.P. wrote the manuscript with substantial feedback from all co-authors.

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