Valence, exchange interaction, and location of Mn ions in polycrystalline Mn$_x$Ga$_{1-x}$N ($x \leq 0.04$)


1Laboratory for Neutron Scattering, Paul Scherrer Institut, CH-5232 Villigen PSI, Switzerland
2Department of Chemistry and Biochemistry, University of Bern, CH-3012 Bern, Switzerland
3Neutron Scattering Division, Oak Ridge National Laboratory, Oak Ridge, Tennessee 37831, USA
4Paul Scherrer Institut, CH-5232 Villigen PSI, Switzerland

(Received 18 October 2017; revised manuscript received 11 January 2018; published 10 April 2018)

We present an experimental study for polycrystalline samples of the diluted magnetic semiconductor Mn$_x$Ga$_{1-x}$N ($x \leq 0.04$) in order to address some of the existing controversial issues. X-ray and neutron diffraction, x-ray absorption near-edge structure, and electron paramagnetic resonance experiments were used to characterize the structural, electronic, and magnetic properties of the samples, and inelastic neutron scattering was employed to determine the magnetic excitations associated with Mn monomers and dimers. Our main conclusions are as follows: (i) The valence of the Mn ions is 2+. (ii) The Mn$^{2+}$ ions experience a substantial single-ion axial anisotropy with parameter $D = 0.027(3)$ meV. (iii) Nearest-neighbor Mn$^{2+}$ ions are coupled antiferromagnetically. The exchange parameter $J = -0.140(7)$ meV is independent of the Mn content $x$; i.e., there is no evidence for hole-induced modifications of $J$ towards a potentially high Curie temperature postulated in the literature.

DOI: 10.1103/PhysRevB.97.140102

There is an ongoing debate on the physical properties of the compound Mn$_x$Ga$_{1-x}$N, a diluted magnetic semiconductor (DMS) with potential applications in spintronics and blue-light emitting diode (LED) technologies [1,2]. The interest in Mn$_x$Ga$_{1-x}$N is driven by the prediction of Mn-induced ferromagnetism with Curie temperatures $T_C$ exceeding room temperature [3], which is required for a technological breakthrough in the field of DMSs. Up to the present, a large number of experiments have been performed for Mn$_x$Ga$_{1-x}$N compounds, but the conclusions still remain highly controversial as recently summarized by Nelson et al. [4]. A basic problem with Mn$_x$Ga$_{1-x}$N is the low solubility of Mn ions in the host compound GaN, so that the investigated samples are often contaminated by Mn clusters or other phases which are ferromagnetic in nature, e.g., Mn$_x$Ga$_{1-x}$Ga ($T_C \approx 770$ K) [5] and Mn$_x$N (ferrimagnetic, $T_C = 738$ K) [6]. For this reason, the observation of ferromagnetism above room temperature reported in the literature has to be considered with caution, especially as none of these findings has resulted in a device working at room temperature. There are some other important questions associated with Mn$_x$Ga$_{1-x}$N for which, so far, no common agreement has been obtained, namely: (i) What is the valence of the Mn ions (Mn$^{2+}$ vs Mn$^{3+}$)? (ii) What are the nature and the size of the magnetic exchange interaction (ferromagnetic vs antiferromagnetic)? (iii) Where are the Mn ions located (regular Ga positions vs interstitial positions)? For an overview of all these aspects we refer to Refs. [13–21] (exchange) and [34–51] (valence) cited by Nelson et al. [4] as well as to the review articles by Liu et al. [7], Jungwirth et al. [8], and Sato et al. [9].

It is the purpose of the present work to provide answers to all these questions through different experiments on polycrystalline samples of Mn$_x$Ga$_{1-x}$N. We describe in the first part the detailed characterization of the samples, which is an indispensable issue, since some of the existing controversies are due to the lack of information on the sample properties. We continue by presenting inelastic neutron scattering (INS) investigations of the magnetic excitations associated with Mn monomers and dimers for which, so far, no information is available in the literature. As a result of the different experimental techniques applied to Mn$_x$Ga$_{1-x}$N, we arrive at consistent conclusions as summarized in the abstract.

Based on the procedure outlined by Szyszkowski et al. [10], polycrystalline Mn$_x$Ga$_{1-x}$N samples were synthesized with manganese concentrations up to $x = 0.08$. The Mn content $x$ was determined by energy dispersive x-ray (EDX) spectroscopy. Analyses by x-ray and neutron diffraction showed that Mn$_x$Ga$_{1-x}$N crystallizes in the hexagonal space group $P6_3mc$, but even for low values of $x$ an impurity phase of type MnN$_y$ ($y < 1$) with tetragonal space group $I4/mmm$ was always present. We concentrated our experimental study on detailed investigations of two samples with $x_1 = 0.024(3)$ and $x_2 = 0.072(2)$ from EDX. The fractional weight of the impurity phase determined by x-ray diffraction amounted to 0.43(3) and 3.7(5)% respectively. Similar values 0.38(7)% and 2.95(15)%, respectively, were obtained from neutron diffraction. Thus, the actual Mn content $x$ of the main phase Mn$_x$Ga$_{1-x}$N has to be corrected accordingly, i.e., for $x_1 = 0.024(3)$ and $x_2 = 0.072(2)$ we have Mn$_{0.02}Ga_{0.98}N$ and Mn$_{0.01}Ga_{0.99}N$, with typically 10% uncertainty for $x$.

The neutron powder diffraction experiments were performed with use of the high-resolution diffractometer for thermal neutrons, HRPT ($\lambda = 1.155$ Å, high-resolution mode with $2d/d = 10^{-3}$) [11], at the spallation neutron source SINQ at PSI Villigen. The refinements of the crystal structures were carried out with the program FULLPROF [12]. The diffraction data taken at $T = 293$ K were analyzed both for the main phase Mn$_x$Ga$_{1-x}$N and for the impurity phase MnN$_y$. In model I the
TABLE I. Neutron diffraction results obtained at room temperature for Mn$_{90}$Ga$_{90}$N in the structure model $P6_3/mmc$ (No. 186) with lattice parameters $a$, $b$, and $c$. Ga is in the (2) position (2/3, 1/3, 0), and N and Mn in the (2b) position (2/3, 1/3, z). $B$ denotes the isotropic displacement factor and $p$ the occupation number. The reliability factors $R_p$ and $\chi^2$ are defined in Ref. [12]. The following constraints were applied: $B$(Ga) = $B$(Mn); $p$(Ga) + $p$(Mn) = 1; $p$(N) = 1. $z$(Mn) was kept fixed for model III at the value obtained from the Fourier analysis. For all models the impurity phase Mn$_N$ was refined in the space group $I4/mmm$ with $y = 0.83$ and resulting lattice parameters $a = b = 2.9762(5)$ Å and $c = 4.1311(13)$ Å.

<table>
<thead>
<tr>
<th></th>
<th>Model I</th>
<th>Model II</th>
<th>Model III</th>
</tr>
</thead>
<tbody>
<tr>
<td>$a (=b)$ (Å)</td>
<td>3.19006(3)</td>
<td>3.19011(3)</td>
<td>3.19007(3)</td>
</tr>
<tr>
<td>$c$ (Å)</td>
<td>5.18608(5)</td>
<td>5.18614(5)</td>
<td>5.18608(5)</td>
</tr>
<tr>
<td>$z$(N)</td>
<td>0.37790(10)</td>
<td>0.37716(13)</td>
<td>0.37782(10)</td>
</tr>
<tr>
<td>$z$(Mn)</td>
<td>0</td>
<td>0.0597(44)</td>
<td>0.18</td>
</tr>
<tr>
<td>$B$(N) ($\tilde{A}^2$)</td>
<td>0.327(16)</td>
<td>0.460(16)</td>
<td>0.352(10)</td>
</tr>
<tr>
<td>$B$(Ga) ($\tilde{A}^2$)</td>
<td>0.321(23)</td>
<td>0.175(19)</td>
<td>0.281(13)</td>
</tr>
<tr>
<td>$B$(Mn) ($\tilde{A}^2$)</td>
<td>0.321(23)</td>
<td>0.175(19)</td>
<td>0.281(13)</td>
</tr>
<tr>
<td>$p$(Ga)</td>
<td>1.012(8)</td>
<td>0.945(4)</td>
<td>0.981(2)</td>
</tr>
<tr>
<td>$p$(Mn)</td>
<td>-0.012(8)</td>
<td>0.055(4)</td>
<td>0.019(2)</td>
</tr>
<tr>
<td>$R_p$</td>
<td>3.15</td>
<td>3.10</td>
<td>3.13</td>
</tr>
<tr>
<td>$R_{wp}$</td>
<td>3.96</td>
<td>3.88</td>
<td>3.93</td>
</tr>
<tr>
<td>$R_{exp}$</td>
<td>2.54</td>
<td>2.54</td>
<td>2.54</td>
</tr>
<tr>
<td>$\chi^2$</td>
<td>2.43</td>
<td>2.32</td>
<td>2.39</td>
</tr>
</tbody>
</table>

FIG. 1. Room temperature XANES spectra measured around the Mn K edge ($\approx 6.555$ keV) for Mn$_{10}$Ga$_{90}$N ($x \approx 0.02$ and $x \approx 0.04$) as well as for the reference samples MnO (Mn$^{2+}$) and Mn$_2$O$_3$ (Mn$^{3+}$).

sites (2/3, 1/3, 0) is identical with tetrahedral coordination, which is an important aspect for the analysis of both the x-ray absorption near-edge structure (XANES) and the INS experiments described below.

XANES experiments at the Mn K edge (6.539 keV) were performed at room temperature with use of the instrument SuperXAS [14] at the Swiss Light Source (SLS) at PSI Villigen. A 2.9-T superconductor magnet provided x rays, and a Si collimating mirror at 2.5 mrad rejected the higher harmonics. A channel-cut Si(111) monochromator was used to select the desired photon energy. A Rh-coated toroidal mirror focused the beam to 0.5 and 0.1 mm in horizontal and vertical dimensions, respectively. For the calibration of the beam energy we used an Fe foil (Fe K edge at 7.112 keV). Mn$_2$O$_3$ and MnO powders served as reference samples for Mn$^{2+}$ and Mn$^{3+}$ in octahedral coordination, respectively. All the samples were measured in the transmission mode. Figure 1 shows the normalized Mn K-edge XANES data recorded for the four samples, which can be interpreted as follows: (i) For Mn$_{10}$Ga$_{90}$N with $x = 0.02$ the energy of the K edge is very close to that of MnO, which favors Mn$^{2+}$. (ii) Raising the Mn content to $x = 0.04$ slightly increases the oxidation state of the Mn ions (probably due to the increased weight of the impurity phase Mn$_N$), but it still remains close to $2^+$. (iii) The strong preedge features around 6.54 keV suggest that the Mn ions have tetrahedral coordination [15], in contrast to the reference samples Mn$_2$O$_3$ and MnO with octahedral Mn coordination. Similar results were obtained from XANES experiments performed for Mn$_{10}$Ga$_{90}$N (0.03 < $x$ < 0.09) layers grown by molecular beam epitaxy on [0001] SiC substrates [16].

Electron paramagnetic resonance (EPR) experiments performed with a microwave frequency of 9.39 GHz gave rise to a resonance at a magnetic field of 0.33 T, similar to the results described by Zajac et al. [17], which agrees with the field-induced splitting of the $m = \pm 1/2$ ground state of Mn$^{2+}$ ions for $g = 2$. In addition, analyses of magnetic susceptibility and magnetization data also favor Mn$^{2+}$.

INS experiments were carried out with the use of the high-resolution time-of-flight spectrometer CNCS [18] at the

dopant Mn ions were treated as substitutional ions at the Ga site with position (2/3, 1/3, 0). However, the refinement provided an occupation number of Mn close to zero. Consequently, in further data refinements we allowed the $z$ coordinate of the Mn position (1/3, 2/3, z) to be varied, resulting in a $z$ displacement of about 0.3 Å from the regular Ga site for model II. Model III was triggered by the results of a Fourier analysis, based on the difference between the observed and the calculated structure factors, which gave evidence for additional scattering from Mn at the interstitial site (2/3, 1/3, z) with $z = 0.18(3)$. The structure parameters obtained for the three models are summarized in Table I.

Models I and III are based on fixed $z$ coordinates of the Mn ions. If we start the refinement with the initial parameters of models I and III by releasing the $z$ constraint of the Mn position, the fitting procedure converges to the parameters of model II. We therefore feel that model II provides the most probable solution for the structure of Mn$_{10}$Ga$_{90}$N. Another argument for model II (and against model III) is the fact that in principle there is no room for placing ions at interstitial sites in the densely packed Wurtzite crystal structure of Mn$_{10}$Ga$_{90}$N, unless either N or Ga vacancies are created in order to comply with the interatomic distances. This is in contrast to the related DMS Mn$_{10}$Ga$_{90}$As, where the dopant Mn ions can occupy interstitial sites commensurate with the zinc-blende crystal structure [13]. Due to the small differences of the reliability factors listed in Table I for the three models, there remains some uncertainty concerning the $z$ coordinate of the Mn ions. However, it is important to realize that the local symmetry of Mn ions being either at the sites (2/3, 1/3, z) or at regular Ga
spallation neutron source (SNS) at Oak Ridge National Laboratory. The samples were enclosed in aluminum cylinders of 8 mm diameter and placed into a He cryostat to achieve temperatures \( T \geq 1.7 \) K. Additional experiments were performed for vanadium to allow the correction of the raw data with respect to background, detector efficiency, and absorption according to standard procedures. We searched for magnetic excitations over a wide energy range, but we did not observe magnetic intensity for energy transfers \( >0.6 \) meV. Energy spectra taken for \( \text{Mn}_{x}\text{Ga}_{1-x}N \) with \( x = 0.04 \) in the neutron energy-gain configuration are shown in Fig. 2(a) for moduli of the scattering vector \( Q \) in the range \( 0.5 \leq Q \leq 1.5 \) \( \text{Å}^{-1} \). The energy of the incoming neutrons was 1.55 meV, with instrumental energy resolutions of Gaussian shape increasing from 20 to 29 \( \mu \)eV for energy transfers from 0 to 0.5 meV, respectively. We observe an increase of the intensity upon raising the temperature from 1.7 to 6.0 K. This becomes more clear by plotting the difference of the energy spectra as shown in Fig. 2(b), which has the advantage that uncertainties about the background are automatically eliminated. Taking intensity differences has proven to be an extremely powerful procedure to analyze INS data [19].

The data of Fig. 2(b) exhibit three partially resolved lines, which were analyzed by Gaussians without any constraints in the least-squares fitting procedure, except for fixing the background at zero intensity. The results are shown as full and broken curves in Fig. 2(b). We interpret the three lines in terms of Mn multimer transitions associated with \( \text{Mn}_{x}\text{Ga}_{1-x}N \). We can neglect the scattering contributions from the impurity phase \( \text{MnN} \). \( \text{MnN} \) and \( \text{Mn}_{3}\text{N}_{2} \) order antiferromagnetically below very high Néel temperatures \( T_N = 660 \) K and \( T_N = 920 \) K, respectively, with nearest-neighbor exchange parameters of the order of \(-20 \) meV [20], giving rise to a spin-wave density of states far above the energy window covered by the present INS experiments.

For the low Mn content only monomers and dimers have to be considered, which based on a random distribution of \( x \) Mn ions over the positions \((1/3, 2/3, z)\) occur with probabilities \( p_{M} = (1-x)^{12} \) and \( p_{D} = 6x(1-x)^{18} \), respectively. The linewidths are considerably enhanced beyond the instrumental energy resolution due to local structural effects [21]. A proper identification of the lines is possible by considering both the spin Hamiltonian and the neutron cross section for monomers and dimers. The spin Hamiltonian of Mn monomers is given by

\[
H = D(s^z)^2, \tag{1}
\]

where \( D \) is the axial single-ion anisotropy parameter, and \( s^z \) denotes the \( z \) component of the spin operator \( s \) of the Mn ions. The corresponding neutron cross section for monomer transitions \( |m\rangle \rightarrow |m'\rangle \) is defined by [22]

\[
\frac{d^2\sigma}{d\Omega d\omega} \propto F^2(Q) n_m(T) |T_1^{\Delta m}|^2, \tag{2}
\]

where \( F(Q) \) is the magnetic form factor, \( n_m(T) \) the Boltzmann population factor of the initial state \( |m\rangle \), and \( -s^z \leq m \leq s^z \). The transition matrix element \( |T_1^{\Delta m}| \) gives rise to the dipole selection rules [22],

\[
\Delta m = m - m’ = 0, \pm 1. \tag{3}
\]

The spin Hamiltonian of Mn dimers is given by

\[
H = -2J_1 \mathbf{s}_1 \cdot \mathbf{s}_2 + D \left[ (s_1^z)^2 + (s_2^z)^2 \right] \tag{4}
\]

where \( J \) is the bilinear exchange parameter. It is convenient to base the diagonalization of Eq. (4) on the dimer states \( |S, M\rangle \), where \( \mathbf{S} = \mathbf{s}_1 + \mathbf{s}_2 \) is the total spin and \(-S \leq M \leq S \). For \( \text{Mn}^{2+} \) ions with \( s_j = 5/2 \) (\( j = 2 \), ferromagnetic \( J > 0 \) and antiferromagnetic \( J < 0 \) exchange give rise to an \( S = 5 \) \((S = 4) \) and an \( S = 0 \) \((S = 0) \) ground state, respectively. The anisotropy term has the effect of splitting the spin states \( |S\rangle \) into the substates \( |S, \pm M\rangle \). The low-energy level schemes for \( \text{Mn}^{2+} \) and \( \text{Mn}^{3+} \) monomers and antiferromagnetically coupled dimers are illustrated in Fig. 3.

![Figure 2](image-url)

**FIG. 2.** (a) Energy spectra of neutrons scattered from \( \text{Mn}_{0.04}\text{Ga}_{0.96}N \) \((x = 0.04)\) in the neutron energy-gain configuration at \( T = 1.7 \) K and \( T = 6.0 \) K. The error bars have the size of the symbols. The incoming neutron energy was 1.55 meV. (b) Difference energy spectrum \( I(T = 6.0 \text{K}) - I(T = 1.7 \text{K}) \) for \( \text{Mn}_{x}\text{Ga}_{1-x}N \) \((x = 0.04)\). (c) Difference energy spectrum \( I(T = 6.0 \text{K}) - I(T = 1.7 \text{K}) \) for \( \text{Mn}_{x}\text{Ga}_{1-x}N \) \((x = 0.02)\). The lines correspond to Gaussian least-squares fits as described in the text. The arrows mark the observed transitions.
The neutron cross section for dimer transitions \(|S,M\rangle \rightarrow |S',M'\rangle\) is defined by [23]

\[
\frac{d^2\sigma}{d\Omega d\omega} \big|_{\Delta M=0} \propto F^2(Q)m_{|S,M\rangle}(T) \left\{ \frac{2}{3} + (-1)^{\Delta S} \left[ \frac{2 \sin(QR)}{(QR)^3} - \frac{2 \cos(QR)}{(QR)^2} \right] \right\} |T_{1\Delta M=0}|^2,
\]

\[
\frac{d^2\sigma}{d\Omega d\omega} \big|_{\Delta M=\pm 1} \propto F^2(Q)m_{|S,M\rangle}(T) \left\{ \frac{2}{3} - (-1)^{\Delta S} \left[ \frac{2 \sin(QR)}{(QR)^3} - \frac{2 \cos(QR)}{(QR)^2} - \frac{\sin(QR)}{QR} \right] \right\} |T_{1\Delta M=\pm 1}|^2,
\]

where \( R \) is the distance between the two dimer spins. The transition matrix element \(|T_{1\Delta M}|\) carries essential information to derive the selection rules through the symmetry properties of the built-in 3- and 6- \( j \) symbols [23]:

\[
\Delta S = S - S' = 0, \pm 1; \quad \Delta M = M - M' = 0, \pm 1. \quad (6)
\]

The \( Q \) dependence of the intensities of the observed multimer transitions displayed in Fig. 4 allows a proper peak identification. The intensity of the line \( M_1 \) follows the form-factor behavior for monomers described by Eq. (2), whereas the intensities of the lines \( D_1 \) and \( D_2 \) are governed by Eq. (5) for dimer transitions. We analyzed the dimer lines \( D_1 \) and \( D_2 \) in Fig. 2(b) on the basis of Eqs. (4)–(6) for both \( \text{Mn}^{2+} \) and \( \text{Mn}^{3+} \) ions. In order to cover all possible values of the ratio \( D/J \), we introduce a parametrization scheme by putting \( D = W_y \) and \( J = W(1 - |y|) \), where \( W \) is an energy-scale factor and \( -1 \leq y \leq 1 \). It follows that \( D/J = 0 \) for \( y = 0 \), while \( D/J = \pm \infty \) for \( y = \pm 1 \). Out of a complete search for \(-1 \leq y \leq 1 \) (with steps \( \Delta y = 0.1 \)), agreement between the observed and calculated data could only be obtained for the parameter sets

\[
\text{Mn}^{3+} : D = 0.108(5) \text{meV}, \quad J = -0.139(10) \text{meV}. \quad (7)
\]

\[
\text{Mn}^{2+} : D = 0.024(3) \text{meV}, \quad J = -0.140(7) \text{meV}. \quad (8)
\]

That is, the Mn dimers are antiferromagnetically coupled. Similar values \( J = -0.16(3) \text{meV} \) [24] and \( J = -0.136 \text{meV} \) [25] were derived from magnetization and magnetic susceptibility data, respectively.

From the analysis of the dimer transitions both \( \text{Mn}^{2+} \) and \( \text{Mn}^{3+} \) are possible; thus we have to consider also the monomer transitions. The allowed monomer transitions are indicated in Fig. 3 by the arrows \( M_1 \) and \( M_2 \). Since the thermal populations of the first-excited monomer states (\(|\pm 3/2\rangle\) for \( \text{Mn}^{2+} \) and \(|\pm 1\rangle\) for \( \text{Mn}^{3+} \)) are practically equal at \( T = 1.7 \text{K} \) and \( T = 6.0 \text{K} \), the transition \( M_2 \) cannot be observed in the difference spectrum, so that only the monomer transition \( M_1 \) is accessible. According to Fig. 3, the monomer transition \( M_1 \) shown in Fig. 2(b) clearly has to be associated with \( \text{Mn}^{2+} \). The resulting single-ion anisotropy parameter calculated from Eq. (1) is \( D = 0.029(3) \text{meV} \), in reasonable agreement with the value derived from the dimer transitions; see Eq. (7).

INS experiments were also performed for \( \text{Mn}_x \text{Ga}_{1-x} \text{N} \) with \( x = 0.02 \) as shown in Fig. 2(c). The resulting energy spectra turned out to be very similar to those for \( x = 0.04 \) displayed.
is too low to generate a sizable ferromagnetic component to the exchange coupling through a hole-mediated mechanism such as Zener’s kinetic exchange interaction [26]. In fact, the concentration of mobile holes was measured to be \( <10^{18} \text{ cm}^{-3} \) for crystalline Mn\(_x\)Ga\(_{1-x}\)N \((x<0.1)\) [24]. A similar number was reported for the DMS compound Mn\(_x\)Zn\(_{1-x}\)Te \((x \leq 0.05)\) where the exchange coupling determined by INS experiments experienced a marginal shift of not more than 1% due to the hole-mediated interaction [27].

The analysis of the neutron diffraction data did not provide a definite answer concerning the location of the dopant Mn\(^{2+}\) ions in Mn\(_x\)Ga\(_{1-x}\)N, although model II (see Table I) is favored for reasons given above. In order to arrive at a conclusive solution of the structure, investigations on single crystals or further neutron diffraction experiments extended to a larger \(Q\) range (by using neutrons with wavelength of typically 0.5 Å) are highly desirable, which are expected to provide an improved discrimination of the models in terms of the \( \chi^2 \) test.

In conclusion, the realization of ferromagnetic DMSs with high Curie temperature relies on both the large moment of the substituted magnetic ions and carrier-induced ferromagnetic exchange interactions. For the crystalline compound Mn\(_x\)Ga\(_{1-x}\)N we have indeed a large magnetic moment \((s=5/2)\), but our study showed that the nature of the exchange between the Mn\(^{2+}\) ions is antiferromagnetic up to \(x=0.04\). Mn\(_x\)Ga\(_{1-x}\)N samples with larger Mn content \(x\) have been obtained for thin films produced by various techniques such as molecular-beam epitaxy (MBE), metal organic chemical vapor deposition (MOCVD), and ion-assisted deposition (IAD). The latter method was used by Granville et al. [28] to prepare precipitate-free samples of Mn\(_x\)Ga\(_{1-x}\)N with Mn contents up to \(x=0.36\). However, magnetic susceptibility measurements demonstrated that the exchange interaction between the Mn\(^{2+}\) ions remains antiferromagnetic. In view of all these facts we conclude that the ferromagnetism reported in the literature for Mn\(_x\)Ga\(_{1-x}\)N is likely due to the presence of either Mn clusters or ferromagnetic impurity phases. Nevertheless, alternative routes have been proposed to realize ferromagnetic DMSs which are based either on codoping with \(p\)-type elements such as Mg [29] or replacing the Ga\(^{3+}\) ions by Li\(^{1+}\) and Zn\(^{2+}\) ions, so that the substitution of Zn\(^{2+}\) by Mn\(^{2+}\) ions is decoupled from carrier doping. The carrier concentration can then be controlled independently of Mn doping by adjusting the Li-(Mn, Zn) stoichiometry, as demonstrated for Li\(_{1+y}\)(Mn\(_x\)Zn\(_{1-x}\))As \((0.05 \leq y \leq 0.2, 0.02 \leq x \leq 0.15)\) with Curie temperatures up to 50 K [30].

Part of this work was performed at the Swiss Spallation Neutron Source (SINQ) and at the Swiss Light Source (SLS), Paul Scherrer Institut (PSI), Villigen, Switzerland. This research used resources at the Spallation Neutron Source, a DOE Office of Science User Facility operated by the Oak Ridge National Laboratory.
