

# Observation of surface magnons and crystalline electric field shifts in superantiferromagnetic NdCu<sub>2</sub> nanoparticles

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An ensemble of superantiferromagnetic NdCu<sub>2</sub> nanoparticles has been produced to perform a detailed analysis of the magnetic excitations using inelastic neutron scattering. Neutron Diffraction measurements indicate a mean nanoparticle size of  $\langle D \rangle \approx 13$  nm, where the bulk commensurate antiferromagnetic structure is retained at the nanoparticle core. Magnetic measurements evidence the interaction among the magnetic moments located at the nanoparticle surface to be strong enough to establish a spin glass behavior. Specific heat analyses show a broad Schottky contribution, revealing the existence of crystalline electric field. Inelastic neutron scattering analyses disclose that the splitting of the crystalline electric field levels associated with the Nd<sup>3+</sup> ions, as well as the spin-wave excitations that emerged below the Néel transition ( $T_N \approx 6$  K) in polycrystalline NdCu<sub>2</sub> [1] are maintained in the nanoparticle state. We have been able to isolate the scattering contribution arising from the nanoparticle surface where both crystalline electric field splitting and the collective magnetic excitations are well-defined despite the symmetry-breaking. Quantitative analyses of this surface scattering reveal that finite-size effects and microstrain lead to a partial inhibition of the transitions from the ground state to the first excited level, as well as a [positive shift \(~15 %\) of the energy associated to collective magnon excitations](#).

## I. INTRODUCTION

Research on magnetic nanoparticles (MNPs) has already become mature [2, 3] and the ensuing technology transfer, especially in data storage, sensing and biomedicine, has boosted a broad interest in the last two decades [4–7]. Although the influence of the MNP size and/or shape [8] in the magnetic properties have been thoroughly studied in 3d core/shell MNPs [9], some subtleties have remained hidden until recently. These are the effects of the magnetic anisotropy [10] or the magnetic disorder and interparticle coupling in super spin glass (SSG) and superferromagnetic systems [11, 12]. Even more, ferromagnetically-coupled entities, rather than magnetic moments within nanoparticles (the so-called *supraferromagnetic correlations*, [12]) have been revealed in clusters of MNPs [13]. All these manifestations of magnetism are intricately connected to the presence of defects, which are intrinsic to nanocrystalline materials, already considered a new class of disordered solids [14]. The existence of defects, strained lattices and surfaces in MNPs has crucial consequences in the magnetic static arrangements and moment dynamics. This is not surprising, considering that very recent studies underline the role of interfacial zones (boundaries) in which a change of atomic periodicity gives rise to unexpected electric and/or magnetic behaviors [15–17].

In the last decades,  $RX_2$  MNPs (where  $R$  stands for rare-earth) have revealed the advantages to easily tune the magnetic coupling (ferro- or antiferromagnetic) by simply changing  $R$  or the non-magnetic metal ( $X$ )

[18, 19]. When it comes to collective excitations, only a few studies have shown direct evidence of the presence of spin-waves in ensembles of MNPs of hematite and maghemite [20–22] by inelastic neutron scattering (INS), even if the propagation of spin-waves is attracting a lot of attention nowadays [23, 24]. However, little is known about the crystalline electric field (CEF) effects in 4f-MNPs, apart from nanocrystalline pure Tb [25], non-periodic TbCd<sub>6</sub> quasicrystals [26], and multilayers [27]. The influence of particle size, microstrain, and surface effects on the CEF and the magnetic excitations in general remains essentially unexplored.

To fill in this gap, in this work, we have studied the CEF excitations in the paramagnetic state and the magnetic excitations in the magnetically ordered phase of an ensemble of superantiferromagnetic (SAF) MNPs of NdCu<sub>2</sub> by using INS. These ensembles of MNPs consist of small antiferromagnetically ordered entities (cores) that get connected to each other via the spin glass disordered surface, which is a result of the uncompensated AF moments in the vicinity of the surface and taking into account the random anisotropy model [28, 29]. It is worth mentioning that the determination and understanding on how the bulk collective excitations develop at the nanoscale is of basic interest in condensed matter physics. Moreover, it is a must to define the role that the core and the surface/interface part of the MNPs are playing in the moment dynamics. This is at the basis of several problems concerning quantum magnetism (e.g., [30–32]). In addition, the findings for these 4f MNPs will complete those previously studied

in ensembles of Fe-oxides MNPs [20, 33, 34].

## II. EXPERIMENTAL DETAILS

Polycrystalline pellets of NdCu<sub>2</sub> have been produced using an arc furnace (MAM-1, Johanna Otto GmbH) under an Ar atmosphere (99.99%). The alloys were sealed-off under Ar pressure (99.99%) and noscaled via ball milling (high-energy planetary Retsch PM 400/2). We have selected milling times  $t = 2$  and 5 hours, as our previous experience indicates these  $t$  to be enough to achieve NP sizes  $\langle D \rangle$  around 10 nm [18, 35, 36]. Although size dispersion is inherent to the milling route, the aforementioned works have reported size dispersion values that are always less than 15% [37], which is, for the aim of this work, more than enough. In this way, this work aims to track the changes that take place in the nanoparticle state compared to the bulk situation, i.e., not to focus on a particular MNP size. Therefore, a size dispersion does not hamper the main results provided in this work. On the contrary, the use of this grinding technique is crucial to the need to access the sufficient amount of MNPs ( $\sim 12$  g MNPs for each milling time) required for measurements.

The structural characterization has been performed by employing X-ray diffraction (XRD) and neutron diffraction (ND). The former were performed at room temperature in a Bruker D8 Advance diffractometer using Cu-K $\alpha$  ( $\lambda = 1.5418 \text{ \AA}$ ) radiation. The ND patterns were collected at G4.1 instrument (LLB, France) using a wavelength  $\lambda = 2.426 \text{ \AA}$  at several temperatures between  $T = 1.5$  K and 15 K. A measuring time of 8h for each pattern has been used to assure a high signal-to-noise ratio.

The magnetic characterization (static  $M_{DC}(T, \mu_0 H)$  and dynamic  $\chi_{AC}(T, f)$ ) was performed in both Quantum Design QD-PPMS and QD-MPMS (SQUID) magnetometers, within the temperature range of  $T = 2$ –300 K and magnetic fields  $\mu_0 H \leq 5$  T. The dynamic  $\chi_{AC}(T, f)$  behavior was analyzed using an oscillating field  $\mu_0 H = 0.313$  mT and frequencies ( $f$ ) ranging from 0.1 kHz to 10 kHz in the 2–10 K range. No DC-bias field was applied during the measurements.

Heat capacity measurements were performed on compacted disks ( $\sim 5$  mg) following the relaxation method [38] under no external applied field within the temperature range of 2 – 300 K. This technique has allowed us to achieve more information on how the short-range collective excitations, that were already present in the bulk alloy, do survive in the nanoparticle state.

INS measurements were made on 5-h milled NdCu<sub>2</sub>

MNPs, using the neutron time-of-flight spectrometer (IN4, ILL, France) with incident energies of  $E_0 = 8.8$ , 16.7, and 66.7 meV. We have focused on two temperature values,  $T = 10$  and 1.5 K, which, according to the magnetic characterization, correspond to the paramagnetic and magnetic state, respectively. All spectra were corrected for background, absorption, and self-shielding, and normalized to vanadium.

## III. RESULTS AND DISCUSSION

Bulk NdCu<sub>2</sub> crystallizes in the orthorhombic CeCu<sub>2</sub>-type structure with space group  $Imma$  (No. 74). The magnetic ion presents a  $4f$  pancake-like electron cloud and below  $T_N = 6.5$  K, antiferromagnetic order, with complex metamagnetic transitions [39]. The CEF description of polycrystalline bulk NdCu<sub>2</sub> was originally evaluated by Gratz *et al.* [1], providing values for the nine crystalline field parameters  $B_l^m$  ( $l = 2, 4, 6$  and  $m = 0, 2, 4, 6$  with  $m \leq l$ ), with numerical values in the range between  $1.2 \times 10^5$  and 0.134 meV [1]. These are going to be taken as the reference values to check whether they become altered or not in the nanoparticle state. To accomplish this, we will provide first the structural and magnetic characterization, as well as the specific heat  $c_P$  analyses used to access more information regarding short-range correlations. These results are grouped in a section labelled *preliminary characterization*. Then, the inelastic neutron scattering data and analyses are provided to unveil the propagation of collective excitations in the nanoscale.

### A. Preliminary characterization

#### 1. Structural characterization

According to the crystallographic characterization (XRD included in [40]), values of mean MNP size and microstrain are obtained as  $\langle D \rangle = 18.3(1.0)$  nm and  $\eta = 0.62(7)\%$  for  $t = 2$ h and  $\langle D \rangle = 13.0(5)$  nm and  $\eta = 0.59(1)\%$  for  $t = 5$ h-milled NdCu<sub>2</sub> MNPs, respectively. As it can be seen, the  $\eta$  values lie below 1% in all cases, which ensures the good degree of crystallinity of the MNPs, in spite of the existence of a certain degree of microstrain. This distortion affects the whole MNP, although it gets, of course, stronger at the MNP surface, where interfaces are formed.

Fig. 1 includes the measured ND patterns together with the Rietveld refinements (Thompson-Cox-Hastings equations, [41]). There, for the paramagnetic (PM) region [see Figs. 1(a) and 1(c)], the Rietveld refinements give a mean MNP size of  $\langle D \rangle = 16.5(4)$  nm (Bragg factor  $R_B = \sum_i |I_i - I_i^{\text{calc}}| / \sum_i |I_i| = 2.6\%$ ) for 2h-milled MNPs and  $\langle D \rangle = 12.7(1)$  nm ( $R_B = 2.4\%$ ) for 5h-milled MNPs. These values are in good agreement with the

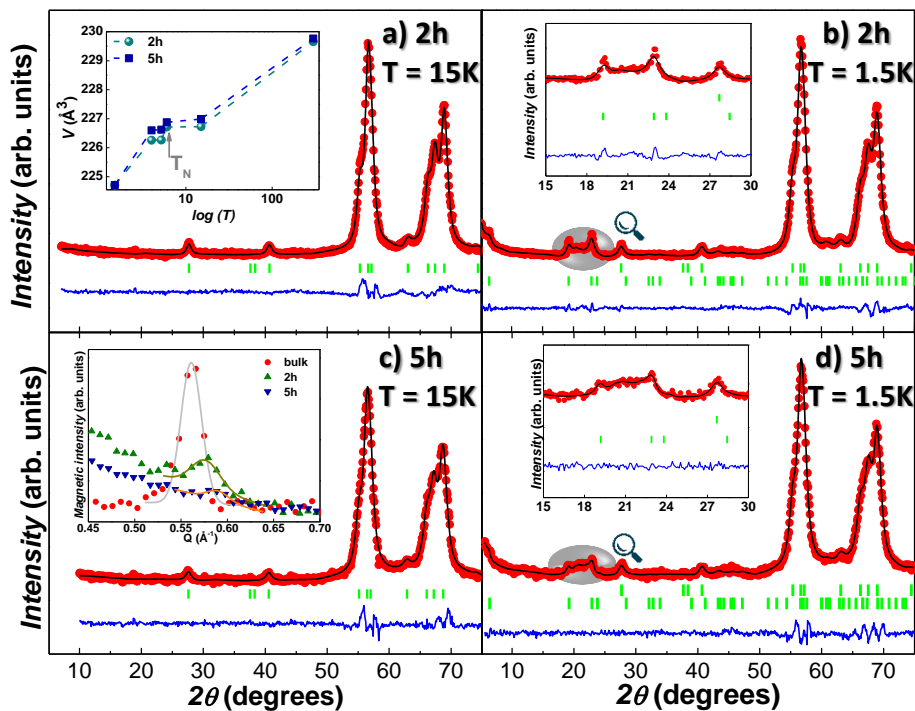


FIG. 1. Neutron diffraction patterns ( $\lambda = 2.426 \text{ \AA}$ ) and Rietveld refinements for (a),(b) NdCu<sub>2</sub> 2h-milled and (c),(d) 5h-milled MNPs measured at (a),(c)  $T = 15 \text{ K}$  and (b),(d)  $T = 1.5 \text{ K}$ , which correspond to the PM and magnetic ordered regions respectively. The inset in (a) shows the reduction of the unit cell volume with decreasing temperature. The position of the Néel transition is marked by the gray arrow. The inset in (c) shows the progressive disappearance of the magnetic peak located at  $Q \approx 0.55 \text{ \AA}^{-1}$  with size reduction. The insets in (b) and (d) allow to observe in closer detail the broadening of the magnetic reflection within the range  $18.5^\circ < 2\theta < 24^\circ$ .

XRD characterization. It is worth noting the occurrence of a magnetoelastic effect, as the unit cell volume  $V$  contraction is affected by the magnetic state of the MNPs. In this way, the decrease rate  $\frac{\Delta V}{\Delta T}$  is  $\approx 42$  times and 33 times greater for  $T < T_N$  than for  $T > T_N$  (PM state). This magnetoelasticity, that is common in RCu<sub>2</sub> intermetallics [1, 42], softens for  $t=5 \text{ h}$  milled MNPs due to the reduction of the number of AF-coupled magnetic moments, driven by the increase of the lattice mismatch [43, 44].

Figures 1(b) and 1(d) concern the magnetic state ( $T = 1.5 \text{ K}$ ). For both ensembles of MNPs, the emergence of magnetic Bragg peaks associated with an AF structure can be observed, especially in the region  $18.5 < 2\theta < 24$  (see the insets). Given that the magnetic unit cell ( $\sim 43 \text{ \AA}$ ) is slightly smaller than the MNP core radius ( $\sim 45 \text{ \AA}$ ), an AF arrangement for the magnetic moments within the core is, then, expected [45]. Both the sharpness and the scattering intensities of these magnetic peaks are reduced with respect to the bulk as a result of finite-size effects. Our magnetic Rietveld refinements indicated that the nuclear size,  $\langle D_N \rangle = 16.5(4) \text{ nm}$  ( $R_B = 2.4\%$ ), whereas the magnetic one is  $\langle D_m \rangle = 15.5(3) \text{ nm}$  ( $R_B = 6.64\%$ ) for 2h-milled MNPs. 5-h milled MNPs display  $\langle D_N \rangle = 12.6(6) \text{ nm}$

( $R_B = 1.56\%$ ) and  $\langle D_m \rangle = 12.4(3) \text{ nm}$  ( $R_B = 4.7\%$ ). The fact that  $\langle D_m \rangle$  is always close to but slightly smaller than  $\langle D_N \rangle$  is due to the presence of a disordered moment arrangement, which is connected to the existence of interfaces. The rise in the magnetic scattering intensity observed in the low- $Q$  region ( $2^\circ < 2\theta < 15^\circ$ , i.e.,  $Q < 0.665 \text{ \AA}^{-1}$ , [see inset of Fig. 1(c)]) is indicative for interparticle correlations, which are triggered by dipolar interactions among MNPs [18, 37]. The reduction in the AF-coupled magnetic moments with size reduction is also supported by the gradual removal of the intense bulk magnetic peak located at  $Q \approx 0.55 \text{ \AA}^{-1}$ , i.e.,  $2\theta \approx 12.4^\circ$ , as this peak gets progressively broaden and displaced to higher  $Q$  values with the size reduction.

The magnetic structure of the MNP cores can be fitted according to a commensurate square-up modulation, in the same way as polycrystalline bulk NdCu<sub>2</sub> [46]. Each magnetic moment, which is oriented along the  $b$ -direction, is ferromagnetically aligned in the  $b$ - $c$  plane, and separated by  $a/2$  with respect to each other. The propagation vectors are  $\tau = (6/10, 0, 0)$  and the harmonic associated with  $3\tau$ . Given that Nd<sup>3+</sup> ions are  $J = 9/2$ , i.e., Kramers-type, a modulation in the intensity is expected [47]. To account for this modulation, we have employed the following Fourier series:

$\mu = \mu_\tau \sin(2\pi\tau R_i/a + \phi_1) + \mu_{3\tau} \sin(6\pi\tau R_i/a + \phi_3)$ , where  $R_i = 0, a/2, a, \dots, 5a$  (sine-wave modulation), which results in magnetic moments of  $\mu_\tau = 2.76(6) \mu_B$  (2h-MNPs) and  $\mu_\tau = 2.64(6) \mu_B$  (5h-MNPs) associated with the first harmonic and  $\mu_{3\tau} = 0.94(3) \mu_B$  and  $\mu_{3\tau} = 0.98(3) \mu_B$  with the third  $3\tau = (0.2, 0, 0)$  ( $R_B = 10.7\%$  for 2h-MNPs and  $R_B = 6.7\%$  for 5h-MNPs). These values give  $M_0 = \pi/4 \cdot \mu_\tau = 2.2(1) \mu_B$  and  $M_0 = \pi/4 \cdot \mu_\tau = 2.0(5) \mu_B$  magnetic moments values for 2h and 5h-MNPs, respectively, and are smaller than the maximum theoretical magnetic moment of  $3.27 \mu_B$ . This finding, which has also been observed previously in the bulk alloy [1], can be understood by CEF influence on the  $\text{Nd}^{3+}$  ions [48]. The decrease of the magnetic moment value with the MNP size confirms the reduction of the AF-coupled entities.

## 2. Magnetic characterization. Dynamic $\chi_{AC}(T, f)$

The magnetic character of the  $\text{NdCu}_2$  MNPs needs to be defined. The static magnetic behavior is presented in the *Static  $M(T, \mu_0 H)$  characterization* section in [49]. The outcome of the analyses is the presence of a disorder state, which may coexist with AF within the MNPs cores. In order to access more information on this disorder state, dynamic susceptibility results are shown hereunder.

Dynamic susceptibility  $\chi_{AC}(T, f)$  measurements corresponding to the in-phase  $\chi'(T, f)$  (top pannel) and out-of-phase  $\chi''(T, f)$  components (bottom) are plotted in Figure 2. Left-side panels ((a) and (b)) correspond to  $\text{NdCu}_2$  2h-milled MNPs and the right-side ones (c) and (d), to 5h-milled ones. A maximum in  $\chi'(T, f)$  occurs at around  $T \approx 5.6$  K for 2h and  $T \approx 5.5$  K for 5h-milled MNPs. This maximum, which evidences dissipation in  $\chi''(T, f)$ , reduces its value and shifts towards higher temperatures when increasing the frequency, which is a characteristic behavior of freezing transitions [19, 50]. As the magnetic moments located within the core retain the bulk AF state (see ND section), there are the magnetic moments located at the MNP surface the ones responsible for this SG-like state. In addition to this SG-cusp, a maximum in  $\chi'(T, f)$  corresponding to the AF Néel transition should be observed at  $T = T_N$ . The fact that we only observe one single broad peak in  $\chi'(T, f)$  is a consequence of the proximity of both  $T_N$  and  $T_f$  values.

In order to obtain more information about the freezing dynamics, we have analyzed the frequency dependence by determining both  $\delta$ -shift parameter,  $\delta = \ln T_f / \log(2\pi f) + k$ , and the critical slowing down law  $\left(\frac{T - T_{f,0}}{T_{f,0}}\right)^{-z\nu}$ , as it is common practice [50–52]. First,  $\delta$ -shift parameter values are 0.0218(7) for 2h and 0.0301(3) for 5h-milled MNPs, respectively. These

values are larger with respect to the ones reported for canonical spin glasses [50], which is not surprising for MNP ensembles, where the concentration of magnetic ions is larger. Nevertheless, the obtained  $\delta$ -shift parameter values are still below the upper limit of SPM ensembles [53], which allow us to ensure the SG nature underlying the freezing dynamics of  $\text{NdCu}_2$ . These values are smaller than the ones corresponding to SAF  $\text{TbCu}_2$  [35] or SAF  $\text{Tb}_{0.5}\text{La}_{0.5}\text{Cu}_2$  MNPs [18], which suggests a more interacting SG ensemble in  $\text{NdCu}_2$  MNPs. Second, the  $f$ -shift of the  $T_f$  values scales up with the aforementioned dynamic critical slowing down law (see insets in Fig. 2 (a) and (c)). Fitting parameters of  $T_{f,0} = 5.36(3)$  K,  $\tau_0 = 9.88(8) \cdot 10^{-11}$  s and  $z\nu = 5.87(3)$  for 2h-milled MNPs and  $T_{f,0} = 5.05(1)$  K,  $\tau_0 = 1.0(1) \cdot 10^{-10}$  s and  $z\nu = 5.05(1)$  for 5h-ones are obtained. These values are close to the ones reported for  $\text{Tb}_{0.5}\text{La}_{0.5}\text{Cu}_2$  SAF MNP ensembles [18, 35] and lie within the expected range for SG freezing transitions [54]. Finally, given the slight increase of  $\delta$ -shift parameter,  $\tau_0$  and  $z\nu$ , together with the slight decrease of the  $T_{f,0}$  as the MNPs get smaller, a decrease in the interaction among the magnetic moments with the size reduction can be stated. This finding has already been observed for SAF  $\text{GdCu}_2$  MNPs [36], where it has been shown how the disappearance of the AF RKKY interactions harms the interactions among the magnetically disordered moments, resulting in a weakened SG state. It may be possible that, if a further reduction of mean size in the nanoparticles is studied (via an increased milling time), the magnetic structure might result in an overall spin glass state, as was discussed in detail in ref. [36]. Finally, no trace for the bulk reorientation temperature ( $T_R \sim 4.5$  K) is found at the nanoparticle state. Bearing in mind both ND and magnetic measurements, a SAF state should be then ascribed for  $t = 2$ h and  $t = 5$ h milled MNPs.

## 3. Specific heat measurements

Heat capacity measurements are a useful tool that provides an initial guide to unveil the magnetic excitations. Following the common practice [1, 55, 56], the specific heat  $c_P$  has been assumed to be formed by the sum of a phononic  $c_{ph}$ , electronic  $c_{el}$  and magnetic  $c_{mag}$  contributions, as:

$$c_P = c_{ph} + c_{el} + c_{mag} \quad (1)$$

the former two contributions are usually grouped in a single term,  $c_{lattice}$ . Given that there exist two different local coordination and symmetry environments within the MNPs, here we propose a more subtle analysis, where we have calculated two different  $c_{lattice}$  contributions, one that gives account for the MNP core and a second one that is responsible for the MNP surface, where the reduced symmetry and enhanced distortion soften the phonon modes [18, 57]. Both contributions

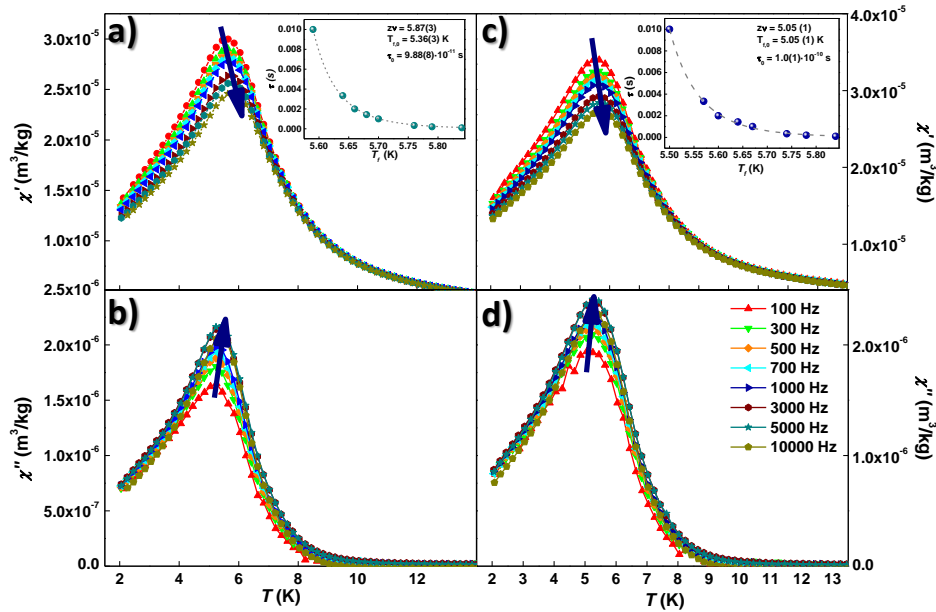


FIG. 2. In-phase  $\chi'(T, f)$  (top panel, (a) and (c)) and out-of-phase  $\chi''(T, f)$  (bottom panel, (b) and (d)) components of the dynamic susceptibility  $\chi_{AC}(T, f)$ , measured applying an oscillating field  $h = 3.13$  Oe, for  $t = 2$  h (left, (a) and (b)) and  $t = 5$  h (right, (c) and (d)) NdCu<sub>2</sub> MNPs. The shift of the maximum associated with the SG state with the frequency of  $h$  is marked with dark blue arrows. Insets exhibit in closer detail the  $\tau \equiv 1/(2\pi f)$  vs.  $T_f$  dependence, where a scaling to a dynamical critical slowing down behavior (power law) is recovered, confirming the freezing (SG) nature of this transition.  $\chi''(T, f)$  component displays dissipation associated with the SG state.

have been weighted, as  $\approx 30\%$  of the magnetic moments are located within the MNP core for a  $\langle D \rangle \approx 13$  nm, assuming a 2 nm-thickness shell (see the *Estimating the core-to-volume ratio* section in [58]). This value for the shell thickness has been chosen based on several stud-

ies performed on diverse ensembles of MNPs, where the surface thickness has been found to be around  $\sim 2$  nm [37, 59, 60]. Accordingly, the  $c_{\text{lattice}}$  contribution has been fitted following:

$$c_{\text{lattice}} = N_c \left[ \gamma_c \cdot T + 9R \left( \frac{T}{\theta_D^c} \right) \int_0^{\theta_D^c/T} dx \frac{x^4 \cdot e^x}{(e^x - 1)^2} \right] + N_s \left[ \gamma_s \cdot T + 9R \left( \frac{T}{\theta_D^s} \right) \int_0^{\theta_D^s/T} dx \frac{x^4 \cdot e^x}{(e^x - 1)^2} \right] \quad (2)$$

where a differentiation between the core and the surface is made. Accordingly, the values for  $\gamma^{\text{core}}$  and  $\theta^{\text{core}}$  have been obtained from the analysis of the  $c_p$  of the bulk alloy, as the magnetic results point to a bulk-like behavior of the magnetic core moments. In this way, we obtain  $\gamma^{\text{core}} = 12.14(13)$  mJ/(molK<sup>2</sup>)<sup>-1</sup> and  $\theta_D^{\text{core}} = 224.7(6)$  K, values that are in good agreement with the ones reported in [1]. On the other hand, we have obtained  $\gamma^{\text{surface}} = 22.82(2)$  mJ/(molK<sup>2</sup>)<sup>-1</sup> and  $\theta_D^{\text{surface}} = 281(4)$  K. These fittings are shown in the right inset of Fig. 3, put into (right panel).

The  $c_{\text{mag}}$  contribution is then obtained following equation Eq. (1) by simply subtracting  $c_{\text{lattice}}$  from the experimental  $c_p$ . Fig. 3 shows the resulting  $c_{\text{mag}}$  (blue spheres), together with the contribution to the specific heat arising from the CEF,  $c_{\text{CEF}}$  (pink line), which has

been calculated as:

$$c_{\text{CEF}} = \frac{1}{kT^2} \cdot \left[ \sum_i E_i^2 p_i - \left( \sum_i E_i p_i \right)^2 \right] \quad (3)$$

being  $p_i = \frac{e^{-E_i/kT}}{\sum_j e^{-E_j/kT}}$  the Boltzmann factor. It should be kept in mind that this contribution gives account for the CEF, thus, it does not include the  $\lambda$ -anomaly arising from the magnetic part.

Beginning from the low temperature region, it is worth noting how the  $\lambda$ -anomaly (zoomed in the green-shaded bottom inset), associated with the core AF state, remains at the same  $T_N = 6.2(1)$  K than the bulk one. Interestingly, no trace for the moment reorientation is observed at  $T_R \sim 4.5$  K which supports the idea of

the weakening of the AF coupling in the MNPs. The reduction of the  $c_{\text{mag}}$  value at  $T_N$  in the MNP state further backs this enfeeblement.

Above  $T_N$ , a broad peak, located between  $10 \text{ K} \lesssim T \lesssim 80 \text{ K}$  is found. This peak, that corresponds to the Schottky anomaly, is caused by the splitting of the energy levels driven by the CEF. This finding is of fundamental interest to the interpretation of collective excitations, as it could be indicating that the CEF level schemes remain almost unaltered in spite of the size reduction and the microstrain of the MNPs. If we compare the experimental  $c_{\text{mag}}$  with the calculated  $c_{\text{CEF}}$ , it can be seen that the temperature value at which the maximum Schottky contribution is found shifts up to higher temperatures for the experimental data, as  $T_{\text{exp}} - T_{\text{calc}} \approx 8 \text{ K}$ . This mismatch between the experimental and the calculated contributions to the specific heat was also detected for the bulk parent alloy, being indicative of the existence of temperature-dependent interactions, such as magnetoelastic effect and/or short-range correlations among  $\text{Nd}^{3+}$ , also reported in [1]. In order to access more information, we have removed these temperature-dependent interactions by normalizing both experimental and calculated contributions to their respective temperature value for which the Schottky anomaly is maximum ( $T/T_{\text{max}}$ ) (see Fig. 3). The measured  $c_{\text{mag}}$  falls to zero above the Schottky anomaly faster than expected according to the calculated  $c_{\text{CEF}}$  (pink line). This indicates that the crystalline electric field  $|9/2\rangle$  multiplet is poorly defined, as its contribution to the specific heat is smooth. This macroscopic insight of the CEF schemes agrees well with the INS results (see below). The possible  $c_{\text{mag}}$  contribution stemming from the SG-coupled moments is too small compared to the one triggered by the CEF (see, for instance, [48], where the canonical  $\text{CuMn}$  SG gives a  $c_{\text{mag}}$  contribution that is 20% the one calculated for the  $\text{NdCu}_2$   $c_{\text{CEF}}$ ).

The dependence of the magnetic entropy  $S_{\text{mag}}$  with the temperature (Fig. 3, top inset) shows that the theoretical value  $S_{\text{mag}}^{\text{theo}}(300 \text{ K}) = R[\ln(2J + 1)] = 19.14 \text{ J/mol}\cdot\text{K}^2$  is not reached, as  $S_{\text{mag}}^{\text{exp}} = 17.1(1) \text{ J/mol}\cdot\text{K}^2$ . The  $S_{\text{mag}}^{\text{exp}}$  is also reduced with respect to the bulk alloy value of  $S_{\text{mag}}^{\text{exp}} = 18.9(1) \text{ J/mol}\cdot\text{K}^2$  (not shown), which is explained by the decrease of the AF-coupled magnetic moments. The fact that four out of five energy levels are populated at  $T \sim 70 \text{ K}$  leads the  $S_{\text{mag}}$  to reach 90% of its maximum value already at this temperature. In the vicinity of  $T_N$ ,  $S_{\text{mag}}^{\text{exp}} \approx 3.85(1) \text{ J/mol}\cdot\text{K}^2$ , which is also below the expected value  $S_{\text{mag}}^{\text{exp}}(T_N) = R[\ln(2)] = 5.76 \text{ J/mol}\cdot\text{K}^2$  for a completely removal of the two-fold moment degeneracy of the CEF ground state doublet. Short-range correlations effects, already argued in polycrystalline  $\text{NdCu}_2$  [1] or  $\text{CeCu}_2$  [61], are at the origin of this finding.

All in all, the  $c_p$  analyses provide initial insight to the

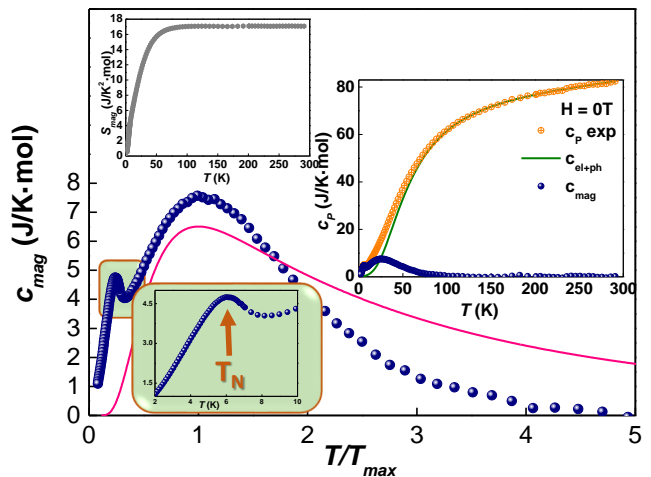


FIG. 3. Magnetic specific heat of  $\text{NdCu}_2$ -5h milled MNPs, where the AF  $\lambda$ -anomaly is found at  $T \sim 6 \text{ K}$  (zoomed in green-bottom inset). This  $c_{\text{mag}}$  contribution has been obtained by subtracting the  $c_{\text{lattice}}$  contribution to the measured  $c_p$  (right inset). The pink line corresponds to the calculated contribution arising from the CEF,  $c_{\text{CEF}}$ . Left top inset corresponds to the  $S_{\text{mag}}^{\text{exp}}(T)$  dependence, where a maximum value of  $S_{\text{mag}}^{\text{exp}} = 17.1(1) \text{ J/mol}\cdot\text{K}^2$  is obtained.

CEF schemes of the  $\text{NdCu}_2$  MNPs, and the presence of the CEF energy level scheme can be anticipated, in spite of the size reduction and the microstrain of the MNPs.

## B. Inelastic Neutron Scattering

The scattering function  $S(Q, \hbar\omega)$  for the 5h-milled  $\text{NdCu}_2$  MNPs is shown in Fig. 4 as a function of wave vector  $Q$  and energy transfer  $\hbar\omega$ . In the PM state at  $T = 10 \text{ K}$ , the excitations corresponding to energy transfer from the ground state to the four excited levels can be distinctly seen for all the measured temperatures in the contours of the scattering function  $S(Q, \hbar\omega)$  [see Figs. 4(d)–4(f)]. The data collected at the higher incident energy  $E_0 = 66.7 \text{ meV}$  [Figs. 4(c) and 4(f)] show that the total splitting  $\sim 13.5 \text{ meV}$  is lower than that of bulk  $\text{NdCu}_2$ ,  $\sim 14.1 \text{ meV}$ . Spectra collected at lower incident energies of 8.8 and 16.7 meV show the different dispersionless excitations from the ground state to the first three excited CEF levels ( $\sim 2.8$ ,  $\sim 5.0$ , and  $\sim 7.3 \text{ meV}$ ). These levels are the same as in bulk  $\text{NdCu}_2$  within experimental precision [1].

In the magnetic state, at  $T = 1.5 \text{ K}$  [Fig. 4(a)–4(c)], the CEF excitations at low energies ( $< 3 \text{ meV}$ ) are strongly modified by the magnetic ordering and the exchange interactions, leading to a dispersive propagating transverse magnetic excitation [1, 62–64], which is seen as a substantially broadened peak due to the powder averaging in our polycrystalline samples. The CEF levels that lie at 4.7 and 7.3 meV in the paramagnetic phase

move up in energy to 5.7 and 7.8 meV, respectively, while higher lying levels are essentially unaffected by the magnetic order.

### 1. CEF level schemes for NdCu<sub>2</sub>

In NdCu<sub>2</sub>, the Nd<sup>3+</sup> ions are located at Wyckoff position 4e with point-group symmetry  $C_{2v}$  and the crystal electric field splits the ten-fold degenerate Hund's rule ground-state multiplet  $^4I_{9/2}$  of the Nd<sup>3+</sup> ions with total angular momentum  $J = 9/2$  into five doublets. The next multiplet state is located several hundreds of meV higher in energy [65]. The CEF and exchange Hamiltonian of  $C_{2v}$  symmetry takes the form:

$$\mathcal{H} = \sum_i \sum_{\ell, m} B_\ell^m O_\ell^m(i) - \frac{1}{2} \sum_{ij} \mathcal{J}_{ij} \mathbf{J}_i \cdot \mathbf{J}_j, \quad (4)$$

where the first term contains the CEF single-ion contributions involving the terms  $\ell = 2, 4, 6$  and  $m = 0, 2, 4$  and  $6$  with  $m \leq \ell$ , where  $O_\ell^m$  and  $B_\ell^m$  are the Stevens operators and CEF parameters, respectively [66]. The second term in Eq. (4) is related to the isotropic RKKY exchange interaction, where  $\mathcal{J}_{ij}$  are the exchange coupling parameters and  $\mathbf{J}_i$  the total angular momentum operator associated with the Nd<sup>3+</sup> ion located on site  $i$ . The magnetic dipole-dipole interactions are not considered.

The CEF and exchange parameters in Eq. (4) were determined from a joint analysis of specific heat, magnetization, magnetic susceptibility, and inelastic neutron-scattering measurements above  $T_N$  [1, 62–64]. These results suggest a doublet CEF ground state formed by  $|0\rangle = 0.049 |\pm 9/2\rangle + 0.891 |\mp 7/2\rangle - 0.373 |\pm 5/2\rangle - 0.230 |\mp 3/2\rangle + 0.111 |\pm 1/2\rangle$  in the basis  $|J, M_J\rangle \equiv |M_J\rangle$ , which is responsible for the uniaxial orthorhombic easy-axis single-ion anisotropy. The first excited state is  $|1\rangle = \mp 0.025 |\pm 9/2\rangle \pm 0.309 |\mp 7/2\rangle \pm 0.306 |\pm 5/2\rangle \pm 0.284 |\mp 3/2\rangle \mp 0.854 |\pm 1/2\rangle$  located at an energy of 2.9 meV and the second excited level is  $|2\rangle = -0.010 |\pm 9/2\rangle + 0.120 |\mp 7/2\rangle + 0.724 |\pm 5/2\rangle - 0.675 |\mp 3/2\rangle + 0.079 |\pm 1/2\rangle$  at 5.0 meV [1]. The remaining two  $J = 9/2$  levels lie at higher energies [see Figs. 4(a)–4(f)]. The latter are retained in the model calculations but are of no relevance for the low-energy magnetic excitations studied in this work or the ground-state properties [1].

Similar to the paradigmatic case of PrNi<sub>2</sub>Si<sub>2</sub>, [67–70], we have used a numerical procedure to calculate the magnetic excitations in the ordered phase within the random phase approximation (RPA), where we treat Eq. (4) in the mean-field approximation [71, 72]. As commented above in Sec. III A 1, the magnetic periodicity is taken into account using a long-period commensurate propagation vector,  $\boldsymbol{\tau} = (6/10, 0, 0) \equiv (0.6, 0, 0)$ . This

means that the magnetic unit cell is assumed to be ten times the length of the body-centered orthorhombic unit cell, corresponding to a period of 20 atomic Nd<sup>3+</sup> layers along the  $a$  axis.

### 2. Neutron Scattering Intensity at the nanoparticle surface

The neutron-scattering intensity is given by:

$$S(\mathbf{Q}, \hbar\omega) = -\frac{1}{\pi} \frac{1}{1 - \exp(-\hbar\omega/k_B T)} |f(\mathbf{Q})|^2 \times \sum_{\alpha\beta} \left( \delta_{\alpha\beta} - \frac{Q_\alpha Q_\beta}{|\mathbf{Q}|^2} \right) \text{Im} \left\{ G_{n=0}^{\alpha\beta}(\mathbf{Q}, \hbar\omega) \right\} \quad (5)$$

where  $Q_\alpha$  are the Cartesian components of the neutron-scattering vector  $\mathbf{Q}$ , and  $f(\mathbf{Q})$  is the magnetic form factor of the Nd<sup>3+</sup> ions. Since the magnetic unit cell has 20 non-equivalent sites,  $n = 0, \dots, 19$ , the corresponding Fourier-transformed two-site Green's functions are coupled within the RPA theory through [71, 73]:

$$\overline{\overline{G}}_N(\mathbf{q}, \hbar\omega) = \overline{g}_N(\hbar\omega) - \sum_{s=0}^{19} \overline{g}_{n-s}(\hbar\omega) \overline{\overline{\mathcal{J}}}(\mathbf{q} + s\boldsymbol{\tau}) \overline{\overline{G}}_s(\mathbf{q}, \hbar\omega), \quad (6)$$

where  $\mathbf{q}$  is the reduced wave vector and  $\overline{g}_N(\hbar\omega)$  is the  $n$ th Fourier component of the single-site dynamic magnetic susceptibility with negative sign calculated from the mean-field levels of the magnetic system shown in Fig. 5. The 20 equations of  $3 \times 3$  matrices in Eq. (6) are solved numerically after the introduction of a small imaginary energy width  $\epsilon$ . The positions of the poles or the excitation energies have also been determined directly using the dynamical matrix-diagonalization method implemented in the McPhase software package [74]. This calculation is used to determine the neutron-scattering intensity  $S_c(\mathbf{Q}, \hbar\omega)$  of the Nd<sup>3+</sup> magnetic moments associated with the core magnetic moments of the MNPs (see below).

Figures 5(a) and 5(b) show  $S(Q, \hbar\omega)$  as a function of energy transfer for a wave vector of  $Q = 1.75 \pm 0.5 \text{ \AA}^{-1}$  at  $T = 10$  and 1.5 K, respectively, obtained by combining data sets taken at different incoming energies. In order to isolate the surface contribution to the total scattering, we have considered the scattering from the MNP state to be a result of the sum of two distinct crystallographic environments, i.e., the nanoparticle core and its surface. Bearing in mind the preliminary characterization shown earlier in this work, we have assumed the magnetic moments located within the core to behave in a way similar to the bulk [39]. The results on the crystalline and magnetic microstructure provided by ND, together with the magnetic characterization, especially,  $\chi_{AC}(T, f)$  dynamic scaling, are unambiguous to this respect. Furthermore, the conservation of the bulk nature at the MNP core was also reported for other RCu<sub>2</sub> MNPs [18, 36, 37]. Given

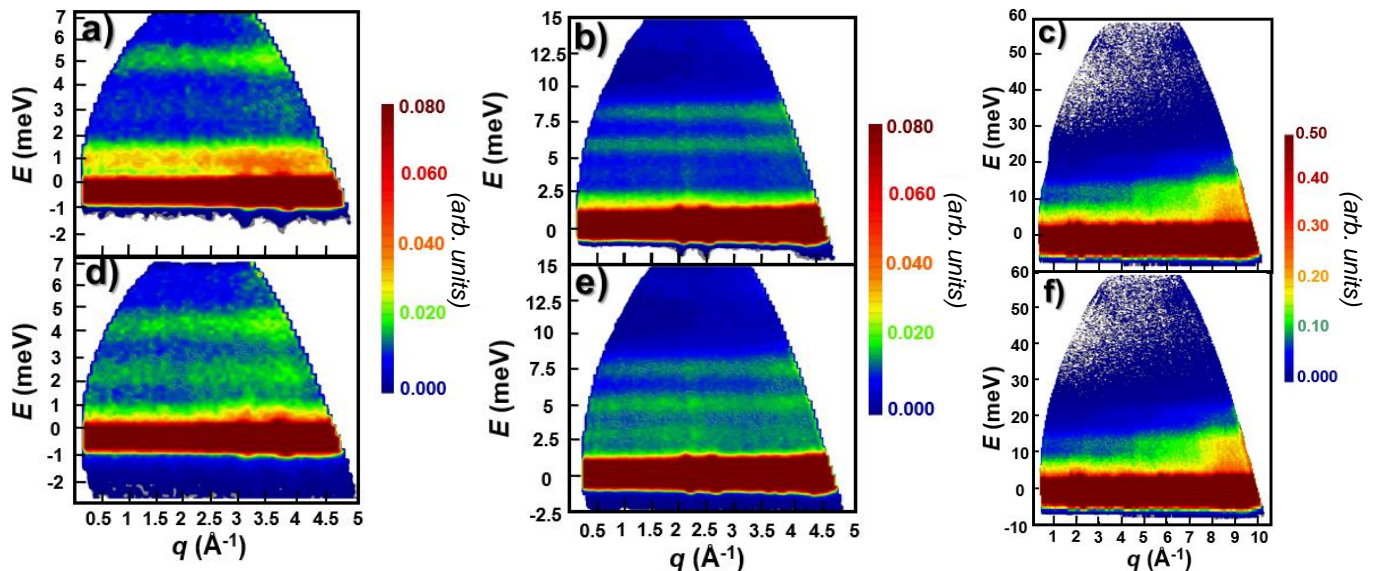


FIG. 4. Contour plots of the scattering function  $S(Q, \hbar\omega)$  as a function of wave vector and energy of 13 nm  $\text{NdCu}_2$  nanoparticles measured at  $T = 1.5$  K in the magnetically ordered phase (top) and at  $T = 10$  K in the paramagnetic phase (bottom), for incoming neutron energies of  $E_0 = 8.8, 16.7,$  and  $66.7$  meV (left to right).

this fact, we can then extract the surface contribution to the INS,  $S_{\text{surface}}(Q, \hbar\omega)$  as:

$$S_{\text{surface}}(Q, \hbar\omega) = S(Q, \hbar\omega) - N S_{\text{core}}(Q, \hbar\omega), \quad (7)$$

where the core contribution  $S_{\text{core}}(Q, \hbar\omega)$  is calculated using the known crystalline–electric–field parameters from the bulk, and  $N$  is a factor that weights the fraction of atoms located within the MNP core. According to the MNP size ( $\langle D \rangle \approx 13$  nm), simple approximations lead to  $N \approx 0.3$ , as the core represents  $\approx 30\%$  of the MNP volume (see the *Estimating the core-to-volume ratio* section in [58]).

The resulting  $S_{\text{surface}}(Q, \hbar\omega)$  is shown in Figs. 5(b) ( $T = 10$  K, PM state) and 5(d) ( $T = 1.5$  K, SAF state). There, both CEF (P1–P4) and collective magnetic excitations can be clearly observed. This is stunning, as it is indicating that the bulk CEF energy level schemes and magnon excitations are well preserved even at the surface of the MNPs. At this stage, it is worth emphasizing that, although we have tried different  $N$  values to give account for this surface  $S_{\text{surface}}(Q, \hbar\omega)$  (see the *Isolating the surface contribution from the total MNP INS intensity* section in [76]), these collective excitations do not disappear for any of the chosen values. Hence, it is reasonable to keep  $N = 0.3$  as it is, according to the aforementioned estimations, the most likely core/MNP ratio for  $\langle D \rangle \approx 13$  nm.

Quantitative information can be obtained by inspecting in closer detail both  $S_{\text{surface}}(Q, \hbar\omega)$  contributions. First, concerning the paramagnetic phase ( $T = 10$  K, Fig. 5 (b)), a left–shift of the energy transfer values can be determined for transitions from the ground state

to the first and to the second excited levels. In this way, CEF excitations are found at  $\hbar\omega_{P1} = 2.6$  and  $\hbar\omega_{P2} = 4.6$  meV for the MNP surface, whereas they were found at  $\hbar\omega_{P1} = 2.9$  and  $\hbar\omega_{P2} = 5.0$  meV for the bulk state [39]. This fact reveals a softening of the CEF splitting, which can be ascribed to the less symmetric crystalline environment in the proximity of the MNP interfaces. This softening may also affect the higher energy excitations (P3 and P4). Nevertheless, the presence of phonon–scattering makes difficult to determine precisely their energy transfer values. Particularly, this scattering becomes rather strong in the energy range from 9 to 14 meV (blue-shadowed region in Fig. 5), which makes specially rough the determination of the P4 position. This phonon–scattering has been confirmed by measuring the INS in non-magnetic isostructural  $\text{YCu}_2$  MNPs (see the *YCu<sub>2</sub>–phonon contribution to INS spectra* section in [75]). Additionally, the poor definition of the  $|9/2\rangle$  multiplet (i.e., P4 excitation) was already foreseen by the specific heat analyses.

On the other hand, change of the relative peak areas between P1 and P2 occurs at the MNP state (Figs. 5 (a) and (b)). In this way, whereas an almost homogeneous ratio between the peak areas  $P2/P1$  was found in bulk  $\text{NdCu}_2$  [1], the value  $P2/P1$  is almost 30% decreased at both the surface and the whole MNP state. This is revealing a partial inhibition of the transitions from the ground state to the first excited level, which reveals that the bulk local–symmetry environment is slightly distorted at the MNPs. The structural microstrain  $\eta$  is at the basis of this distortion. Even if minimal ( $\eta < 1\%$ ), it slightly changes the cation distribution surrounding the  $\text{Nd}^{3+}$  ions, thus, provoking the decoherence of some



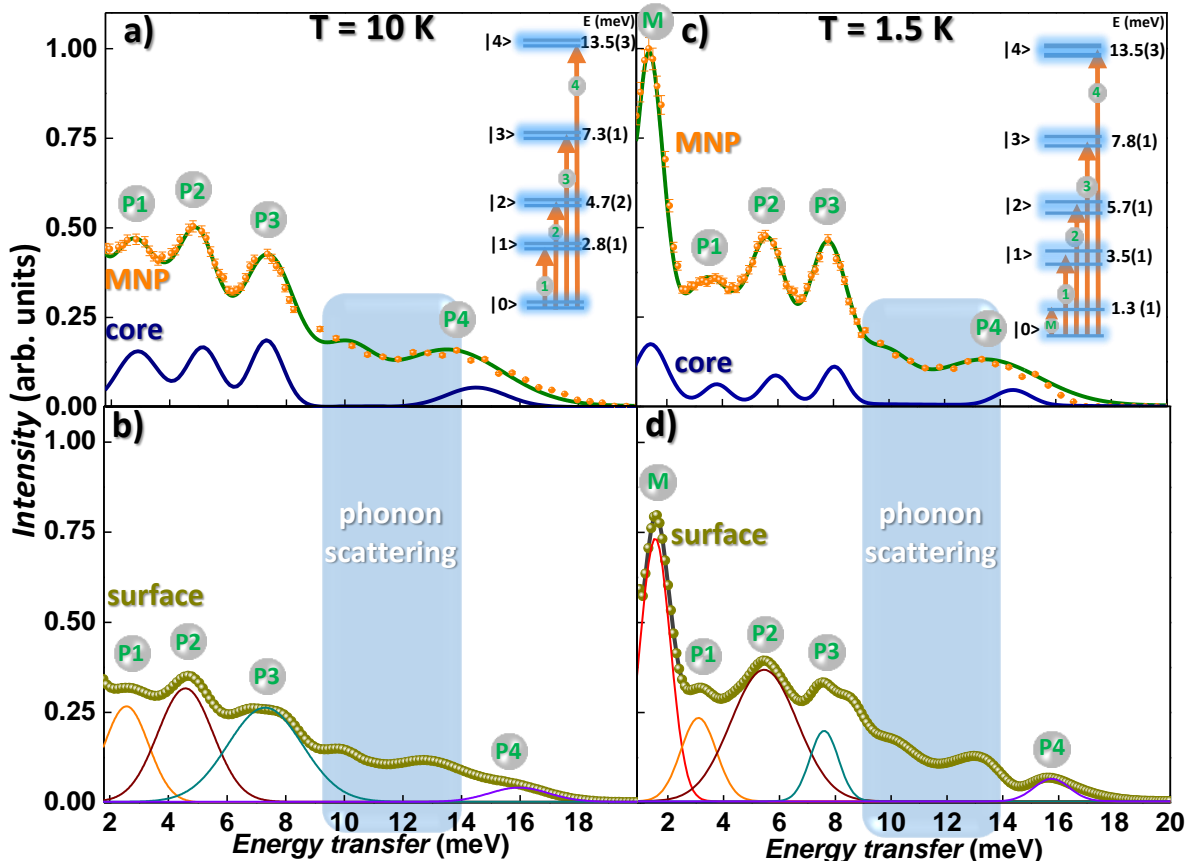


FIG. 5. (a),(c)  $S(Q, \hbar\omega)$  of 5h-milled  $\text{NdCu}_2$  MNPs as a function of energy transfer for a wave vector of  $Q = 1.75 \pm 0.5 \text{ \AA}^{-1}$  (black symbols) at  $T = 10 \text{ K}$  (left) and  $1.5 \text{ K}$  (right), obtained by combining data sets taken at different incoming energies. Error bars for the experimental MNP data are around 3%. Blue lines: core contribution calculated from bulk  $\text{NdCu}_2$  (see text). (b),(d) Surface contribution according to Eq. (7). Thin lines show fits of Gaussians to the data. The blue rectangle ranging from 9 to 14 meV indicates the region where phonon scattering is masking the magnetic signal (see [75]). The insets in (a) and (c) show the CEF level schemes in the paramagnetic and magnetically ordered phases, respectively, where the blue blurred regions around each level represent the distribution of energies.

CEF excitations. Of course, this distortion should also be affecting P2–P4 transitions, but its effect is almost not noticeable within the experimental resolution, given that the distortion is very subtle ( $\eta < 1\%$ ). Therefore, it can only be detected for the most intense excitation (i.e., from the ground state to the first excited level, P1). Given that this partial inhibition from the ground state to the first excited level is close, but slightly greater at the whole MNP  $S(Q, \hbar\omega)$  than the one at the  $S_{\text{surface}}(Q, \hbar\omega)$ , both the core and surface microstrain are definitely playing a role. Therefore, we can state that the intermediate regions between the AF-ordered MNP cores and SG-disordered surfaces in MNPs play the same role as *interfaces* do in multilayers, where a modification of both CEF level splittings and ground-state wave functions was reported (e.g., in Fe–Nd multilayers [27]).

Second, the magnetically ordered phase ( $T = 1.5 \text{ K}$ , Fig. 5(d)) shows the co-existence of collective magnetic excitations and CEF levels, which is actually unprecedented, as the loss of periodicity at interfaces is

well-known to affect the magnetic interactions. Here, the lowest energy excitation (peak labelled M) is found at  $\hbar\omega \approx 1.6 \text{ meV}$ , whereas the bulk one appeared at  $\hbar\omega \approx 1.4 \text{ meV}$  [1]. This represents a shift of  $\sim 15\%$ . Given the minor uncertainty associated with the measured energy transfer  $\hbar\omega$  ( $\sim 3\%$ ), this shift can be considered unambiguous. The investigations carried out by E. Gratz *et al.* [1] on the bulk state related this peak to a transverse spin-wave mode generated from the AF  $\text{NdCu}_2$  ground state. **This positive shift may be ascribed to an enhancement of the anisotropy constant,  $K$ , as it has been shown, for instance, in [77]. In the case of our MNPs, this enhancement could be ascribed to both size-effects and microstrain.**

#### IV. DISCUSSION

We have been able to produce a large quantity ( $\sim 12$

g) of NdCu<sub>2</sub> MNPs by the ball milling route. Relatively low milling times ( $t = 5$  h) have been enough to achieve 13 nm-sized MNPs, in the same way as other RCu<sub>2</sub> MNPs [18, 35–37]. The produced MNPs retained the AF bulk structure at the core and a disordered SG state for the magnetic moments sitting at the surface, which is also common for nano-RCu<sub>2</sub> or RAl<sub>2</sub> [18, 35–37, 78].

The experimental data ( $c_P$ , INS) have shown the occurrence of crystalline electric field and magnon collective excitations for the NdCu<sub>2</sub> 13-nm MNPs. Although the general picture shows that the level schemes remain very similar to the bulk situation, some subtle differences are been disclosed thanks the evaluation of the surface contribution to the total INS, which has been obtained following Eq. (7). This allows to access quantitative information regarding the CEF and magnon collective excitations propagation at the MNP surface. One point that should be addressed concerns the  $N$  value shown in eq. 7. Although it has been estimated in a *simplified* manner (see the section *Estimating the core-to-volume ratio* in [58]), the prevalence of collective excitations at the surface does not dependent on the different  $N$  values. To ensure this, we have considered a wide variety of  $N$  values (see some examples shown in the *Isolating the surface contribution from the total MNP INS intensity* section, [76]) and both the CEF and magnon excitations did remain. In any case, this small uncertainty around the  $N$  value is not a hindrance to the main conclusions of this study.

Regarding the surface INS contribution, the loss of long-range symmetry at the nanoparticle surface has revealed not to be enough to rule out the CEF level schemes. This fact reveals that the CEF is robust against local symmetry distortions. Second, it turns out that transitions to the first excited level at  $T = 10$  K are partially inhibited with respect to the bulk landscape reported in [1]. This partial inhibition is deduced from the decrease in the intensity for transitions from the ground state to the first excited level, and can be attributed to a distortion of the local-symmetry environment surrounding each Nd<sup>3+</sup>-ion. Nevertheless, it should be kept in mind that the distortion of the crystalline structure is below 1%, according to XRD (see [40]). Therefore, this effect is weak, *i.e.*, it is only detected for the first excitation, which is the most intense one. It cannot be discarded, of course, that this distortion could be affecting the rest of the transitions, but as they are less intense, its effect is not noticeable within the experimental resolution. Further INS experiments using different resolutions and/or other MNP ensembles may be helpful to complete this statement.

At  $T = 1.5$  K (magnetic state), a positive shift in the energy associated with the spin-wave excitations is found at the MNP state. [The reason for this displacement \( \$\sim 15\%\$ \) might be ascribed to the enhanced magnetic anisotropy at the nanoparticle state, which](#)

[is supported by the  \$M\(T,H\)\$  characterization \(see the \*Static  \$M\(T,H\)\$  characterization\* section in \[49\]\).](#) Although our experimental evidence is unambiguous, it holds true that this leaves an open path to further INS experiments to be performed, where different MNP sizes could be evaluated. We recall at this point that it is challenging to perform INS in MNP ensembles [20, 21, 33, 34].

## V. CONCLUSION

Inelastic neutron scattering measurements on polycrystalline NdCu<sub>2</sub> nanoparticles of  $\langle D \rangle \approx 13$  nm-size have been successfully carried out. We have extracted the surface contribution from the total scattering intensity, being able to experimentally observe that the crystalline electric field level schemes, as well as magnon excitations, are well-defined at the MNP surface despite the distortion of these outer atoms. In this way, the reduction in the local coordination symmetry, as well as the structural distortion and magnetic disorder of the outer Nd<sup>3+</sup> ions, do not affect significantly the CEF Hamiltonian. Nevertheless, the results and analyses presented in this work have allowed us to deepen inside the subtleties concerning the propagation of collective excitations in the MNP state. In this way, it has been possible to determine a partial inhibition of the transitions from the ground state to the first excited level, as well as a [slight positive shift \( \$\sim 15\%\$ \) of the energy transfer corresponding to magnon excitations](#). Both effects [can be ascribed to finite-size effects](#) (local-symmetry coordination, dimensionality) and microstrain. The present inelastic neutron scattering observations in NdCu<sub>2</sub> pave the way for studies in which interfaces between geometrically ordered and disordered regions are present in the nanocrystalline materials. In particular, it would be interesting to analyze the excitations at higher energy using in similar ensembles of  $4f$  nanoparticles, [as well as to disclose the energy shift of the magnon excitations using a neutron spectrometer with better energy resolution](#).

## VI. ACKNOWLEDGEMENTS

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- [1] E. Gratz, M. Loewenhaupt, M. Divis, W. Steiner, E. Bauer, N. Pillmayr, H. Müller, H. Nowotny, and B. Frick, *J. Phys.: Condens. Matter* **3**, 9297 (1991).
- [2] R. Kodama, *J. Magn. Magn. Mater.* **200**, 359 (1999).
- [3] A.-H. Lu, E. e. Salabas, and F. Schüth, *Angewandte Chemie Int. Ed.* **46**, 1222 (2007).
- [4] G. Reiss and A. Hütten, *Nat. Mater.* **4**, 725 (2005).
- [5] A. Moser, K. Takano, D. T. Margulies, M. Albrecht, Y. Sonobe, Y. Ikeda, S. Sun, and E. E. Fullerton, *J. Phys. D: Appl. Phys.* **35**, R157 (2002).
- [6] L. Gloag, M. Mehdipour, D. Chen, R. D. Tilley, and J. J. Gooding, *Adv. Mater.* **31**, 1904385 (2019).
- [7] Q. A. Pankhurst, J. Connolly, S. K. Jones, and J. Dobson, *J. Phys. D: Appl. Phys.* **36**, R167 (2003).
- [8] P. Guardia, A. Labarta, and X. Batlle, *The J. Phys. Chem. C* **115**, 390 (2011).
- [9] J. Nogués, J. Sort, V. Langlais, V. Skumryev, S. Surinach, J. Muñoz, and M. Baró, *Phys. Rep.* **422**, 65 (2005).
- [10] I. Orue, L. Marcano, P. Bender, A. García-Prieto, S. Valencia, M. Mawass, D. Gil-Cartón, D. A. Venero, D. Honecker, A. García-Arribas, *et al.*, *Nanoscale* **10**, 7407 (2018).
- [11] X. Chen, S. Bedanta, O. Petravic, W. Kleemann, S. Sahoo, S. Cardoso, and P. Freitas, *Phys. Rev. B* **72**, 214436 (2005).
- [12] P. Bender, D. Honecker, and L. Fernández Barquín, *Appl. Phys. Lett.* **115**, 132406 (2019).
- [13] J. Alonso, M. Fdez-Gubieda, J. Barandiarán, A. Svalov, L. F. Barquín, D. A. Venero, and I. Orue, *Phys. Rev. B* **82**, 054406 (2010).
- [14] H. Gleiter, *Prog. Mater. Sci.* **33**, 223 (1989).
- [15] S. Das, Y. Tang, Z. Hong, M. Gonçalves, M. McCarter, C. Klewe, K. Nguyen, F. Gómez-Ortiz, P. Shafer, E. Arenholz, *et al.*, *Nature* **568**, 368 (2019).
- [16] A. Brinkman, M. Huijben, M. Van Zalk, J. Huijben, U. Zeitler, J. Maan, W. G. van der Wiel, G. Rijnders, D. H. Blank, and H. Hilgenkamp, *Nat. Mater.* **6**, 493 (2007).
- [17] D. Zákutná, D. Nižňanský, L. C. Barnsley, E. Babcock, Z. Salhi, A. Feoktystov, D. Honecker, and S. Disch, *Phys. Rev. X* **10**, 031019 (2020).
- [18] E. M. Jefremovas, M. de la Fuente Rodríguez, J. Alonso, J. R. Fernández, J. Espeso, I. Puente-Orench, D. Rojas, A. García-Prieto, M. Fdez-Gubieda, L. R. Fernández, and L. Barquín, *Nanomaterials* **10**, 2148 (2020).
- [19] G. Zhou and H. Bakker, *Phys. Rev. Lett.* **73**, 344 (1994).
- [20] S. N. Klausen, K. Lefmann, P.-A. Lindgård, L. T. Kuhn, C. R. H. Bahl, C. Frandsen, S. Mørup, B. Roessli, N. Cavadini, and C. Niedermayer, *Phys. Rev. B* **70**, 214411 (2004).
- [21] S. Disch, R. P. Hermann, E. Wetterskog, A. A. Podlesnyak, K. An, T. Hyeon, G. Salazar-Alvarez, L. Bergström, and T. Brückel, *Phys. Rev. B* **89**, 064402 (2014).
- [22] J. Robert, E. Lhotel, G. Remenyi, S. Sahling, I. Mirebeau, C. Decorse, B. Canals, and S. Petit, *Phys. Rev. B* **92**, 064425 (2015).
- [23] M. Dabrowski, T. Nakano, D. M. Burn, A. Frisk, D. G. Newman, C. Klewe, Q. Li, M. Yang, P. Shafer, E. Arenholz, *et al.*, *Phys. Rev. Lett.* **124**, 217201 (2020).
- [24] Y. Nambu, J. Barker, Y. Okino, T. Kikkawa, Y. Shiomi, M. Enderle, T. Weber, B. Winn, M. Graves-Brook, J. Tranquada, *et al.*, *Phys. Rev. Lett.* **125**, 027201 (2020).
- [25] S. Philippi, J. Markmann, R. Birringer, and A. Michels, *J. of Appl. Phys.* **105**, 07A701 (2009).
- [26] P. Das, P.-F. Lory, R. Flint, T. Kong, T. Hiroto, S. L. Bud'ko, P. C. Canfield, M. de Boissieu, A. Kreyssig, and A. I. Goldman, *Phys. Rev. B* **95**, 054408 (2017).
- [27] L. Baczewski, M. Piecuch, J. Durand, G. Marchal, and P. Delcroix, *Phys. Rev. B* **40**, 11237 (1989).
- [28] S. Mørup, M. B. Madsen, J. Franck, J. Villadsen, and C. J. Koch, *Journal of Magnetism and Magnetic Materials* **40**, 163 (1983).
- [29] S. Bedanta, T. Eimüller, W. Kleemann, J. Rhensius, F. Stromberg, E. Amaladass, S. Cardoso, and P. Freitas, *Physical Review Letters* **98**, 176601 (2007).
- [30] M. Hermele, M. P. Fisher, and L. Balents, *Physical Review B* **69**, 064404 (2004).
- [31] M. J. Gingras and P. A. McClarty, *Reports on Progress in Physics* **77**, 056501 (2014).
- [32] Y. Shen, Y.-D. Li, H. Wo, Y. Li, S. Shen, B. Pan, Q. Wang, H. Walker, P. Steffens, M. Boehm, *et al.*, *Nature* **540**, 559 (2016).
- [33] M. Hennion, C. Bellouard, I. Mirebeau, J. Dormann, and M. Nogues, *EPL (Europhysics Letters)* **25**, 43 (1994).
- [34] M. F. Hansen, F. Bødker, S. Mørup, K. Lefmann, K. N. Clausen, and P.-A. Lindgård, *Physical review letters* **79**, 4910 (1997).
- [35] C. Echevarria-Bonet, D. P. Rojas, J. I. Espeso, J. R. Fernández, M. de la Fuente Rodríguez, L. F. Barquín, L. R. Fernández, P. Gorria, J. A. Blanco, M. Fdez-Gubieda, E. Bauer, and F. Damay, *J. Phys.: Condens. Matter* **27**, 496002 (2015).
- [36] E. Jefremovas, J. A. Masa, M. Rodríguez, J. R. Fernández, J. Espeso, D. Rojas, A. García-Prieto, M. Fernández-Gubieda, and L. F. Barquín, *Nanomaterials* **10**, 1117 (2020).
- [37] C. Echevarria-Bonet, D. P. Rojas, J. I. Espeso, J. R. Fernández, M. de la Fuente Rodríguez, L. F. Barquín, L. R. Fernández, P. Gorria, J. A. Blanco, M. L. Fdez-Gubieda, E. Bauer, and F. Damay, *Phys. Rev. B* **87**, 180407(R) (2013).
- [38] R. Bachmann, F. DiSalvo Jr, T. Geballe, R. Greene, R. Howard, C. King, H. Kirsch, K. Lee, R. Schwall, H.-U. Thomas, *et al.*, *Rev. of Scientific Instruments* **43**, 205 (1972).
- [39] M. Ellerby, K. McEwen, M. De Podesta, M. Rotter, and E. Gratz, *J. Phys.: Condens. Matter* **7**, 1897 (1995).
- [40] See Supplemental Material at [www.url.com](http://www.url.com) for additional information on X-Ray Diffraction characterization.
- [41] P. Thompson, D. Cox, and J. Hastings, *J Appl. Crystallogr* **20**, 79 (1987).
- [42] P. Morin and D. Schmitt, in *Quadrupolar Interactions and Magneto-Elastic Effects in Rare Earth Intermetallic Compounds* (Elsevier, London, 1990) Chap. 1.
- [43] R. H. Kodama, S. A. Makhlof, and A. E. Berkowitz, *Physical Review Letters* **79**, 1393 (1997).
- [44] N. Rinaldi-Montes, P. Gorria, D. Martínez-Blanco, A. Fertes, L. F. Barquín, J. R. Fernández, I. de Pedro, M. Fdez-Gubieda, J. Alonso, L. Olivi, *et al.*, *Nanoscale* **6**, 457 (2014).

- [45] A. A. Baker, J. R. Lee, C. A. Orme, T. van Buuren, and S. K. McCall, *J. Phys.: Condens. Matter* **32**, 495803 (2020).
- [46] R. Arons, M. Loewenhaupt, T. Reif, and E. Gratz, *J. Phys.: Condens. Matter* **6**, 6789 (1994).
- [47] T. Chatterji, in *Neutron Scattering From Magnetic Materials* (Elsevier, London, 2005) Chap. 6.
- [48] L. Bozukov, A. Gilewski, E. Gratz, A. Apostolov, and K. Kamenov, *Physica B: Condens. Matter* **177**, 299 (1992).
- [49] See Supplemental Material at [www.url.com](http://www.url.com) for additional information on static  $m_{DC}$  characterization.
- [50] J. A. Mydosh, in *Spin Glasses: An Experimental Introduction* (Taylor and Francis, London, 1993) Chap. 4.
- [51] P. Jönsson, M. Hansen, P. Svedlindh, and P. Nordblad, *J. Magn. Magn. Mater.* **226**, 1315 (2001).
- [52] Z. Fu, Y. Zheng, Y. Xiao, S. Bedanta, A. Senyshyn, G. G. Simeoni, Y. Su, U. Rücker, P. Kögerler, and T. Brückel, *Phys. Rev. B* **87**, 214406 (2013).
- [53] S. Bedanta and W. Kleemann, *J. of Phys. D: Appl. Phys.* **42**, 013001 (2009).
- [54] J. Souletie and J. Tholence, *Phys. Rev. B* **32**, 516 (1985).
- [55] M. Bouvier, P. Lethuillier, and D. Schmitt, *Phys. Rev. B* **43**, 13137 (1991).
- [56] N. H. Luong, J. Franse, and T. D. Hien, *J. of Phys. F: Metal Phys.* **15**, 1751 (1985).
- [57] D. Rojas, L. F. Barquín, J. R. Fernández, L. R. Fernández, and J. Gonzalez, *Nanotechnology* **21**, 445702 (2010).
- [58] See Supplemental Material at [www.url.com](http://www.url.com) for sketch and approximations.
- [59] L. Del Bianco, D. Fiorani, A. Testa, E. Bonetti, L. Savini, and S. Signoretti, *Physical Review B* **66**, 174418 (2002).
- [60] Q. A. Pankhurst, Á. Y. Martínez, and L. F. Barquín, *Physical Review B* **69**, 212401 (2004).
- [61] E. Gratz, N. Pillmayr, E. Bauer, H. Muller, B. Barbara, and M. Loewenhaupt, *J. Phys.: Condens. Matter* **2**, 1485 (1990).
- [62] M. Loewenhaupt, T. Reif, W. Hahn, and B. Frick, *J. Magn. Magn. Mater.* **177-181**, 1050 (1998).
- [63] M. Rotter, M. Loewenhaupt, S. Kramp, T. Reif, N. M. Pyka, W. Schmidt, and R. van de Kamp, *Eur. Phys. J. B* **14**, 29 (2000).
- [64] M. Rotter, S. Kramp, M. Loewenhaupt, E. Gratz, W. Schmidt, N. M. Pyka, B. Hennion, and R. van de Kamp, *Appl. Phys. A* **74**, S751 (2002).
- [65] J. H. van Vleck, in *The Theory of Electric and Magnetic Susceptibilities* (Oxford University Press, London, 1932) Chap. 9.
- [66] M. T. Hutchings, *Solid State Phys.* **16**, 227 (1964).
- [67] J. A. Blanco, D. Gignoux, and D. Schmitt, *Phys. Rev. B* **45**, 2529 (1992).
- [68] J. A. Blanco, D. Schmitt, and J. C. Gómez-Sal, *J. Magn. Magn. Mater.* **116**, 128 (1992).
- [69] J. A. Blanco, B. Fåk, E. Ressouche, B. Grenier, M. Rotter, D. Schmitt, J. A. Rodríguez-Velamazán, J. A. Campo, and P. Lejay, *Phys. Rev. B* **82**, 054414 (2010).
- [70] J. A. Blanco, B. Fåk, J. Jensen, M. Rotter, A. Hiess, D. Schmitt, and P. Lejay, *Phys. Rev. B* **87**, 104411 (2013).
- [71] K. A. McEwen, U. Steigenberger, and J. Jensen, *Phys. Rev. B* **43**, 3298 (1991).
- [72] R. M. Nicklow and N. Wakabayashi, *Phys. Rev. B* **51**, 12425 (1995).
- [73] J. Jensen and A. R. Mackintosh, in *Rare Earth Magnetism: Structures and Excitations* (Oxford University Press, Oxford, 1991).
- [74] M. Rotter, M. D. Le, A. T. Boothroyd, and J. A. Blanco, *J. Phys.: Condens. Matter* **24**, 213201 (2012); McPhase software package <http://www.mcphase.de>
- [75] See Supplemental Material at [www.url.com](http://www.url.com) for further information on the phonon–band contribution.
- [76] See Supplemental Material at [www.url.com](http://www.url.com) for various  $s_{\text{surface}}(q, \omega)$  contributions.
- [77] P. Talagala, P. S. Fodor, D. Haddad, R. Naik, L. Wenger, P. Vaishnava, and V. Naik, *Physical Review B* **66**, 144426 (2002).
- [78] D. Rojas, L. F. Barquín, J. S. Marcos, C. Echevarria-Bonet, J. Espeso, J. R. Fernández, L. R. Fernández, and M. Mathon, *Materials Research Express* **2**, 075001 (2015).