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Magnetic correlations in infinite-layer nickelates: An experimental and theoretical multimethod study


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We report a comprehensive study of magnetic correlations in LaNiO_2 , a parent compound of the recently discovered family of infinite-layer (IL) nickelate superconductors, using multiple experimental and theoretical methods. Our specific heat, muon-spin rotation (μSR), and magnetic susceptibility measurements on polycrystalline LaNiO_2 show that long-range magnetic order remains absent down to 2 K. Nevertheless, we detect residual entropy in the low-temperature specific heat, which is compatible with a model fit that includes paramagnon excitations. The μSR and low-field static and dynamic magnetic susceptibility measurements indicate the presence of short-range magnetic correlations and glassy spin dynamics, which we attribute to local oxygen nonstoichiometry in the average infinite-layer crystal structure. This glassy behavior can be suppressed in strong external fields, allowing us to extract the intrinsic paramagnetic susceptibility. Remarkably, we find that the intrinsic susceptibility shows non-Curie-Weiss behavior at high temperatures, in analogy to doped cuprates that possess robust nonlocal spin fluctuations. The distinct temperature dependence of the intrinsic susceptibility of LaNiO_2 can be theoretically understood by a multimethod study of the single-band Hubbard model in which we apply complementary cutting-edge quantum many-body techniques (dynamical mean-field theory, cellular dynamical mean-field theory, and the dynamical vertex approximation) to investigate the influence of both short- and long-ranged correlations. Our results suggest a profound analogy between the magnetic correlations in parent (undoped) IL nickelates and doped cuprates.

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I. INTRODUCTION

Revealing the major mechanism mediating Cooper pairing in unconventional superconductors is one of the defining challenges of condensed matter research [1]. In superconducting cuprates and pnictides, pairing via exchange of magnetic excitations is widely considered to be the most relevant scenario [1–4]. Yet, the nature of the “glue” of the superconductivity in Sr or Ca substituted $R\text{NiO}_2$ ($R = \text{La}, \text{Pr}, \text{Nd}$) [5–10] has not been clarified conclusively [11]. Several theoretical [12–20] and first experimental studies [21–23] on these infinite-layer (IL) nickelates have suggested the presence of cupratelike spin fluctuations, which therefore are a prime candidate for providing the attractive interactions underlying their presumably unconventional superconductivity [17,24,25]. Moreover,

IL nickelates and cuprates share key similarities [26], such as a nominal $3d^9$ electronic configuration of the Ni^{1+} and Cu^{2+} ions with spin $s = \frac{1}{2}$. Furthermore, IL nickelates and IL cuprates are isostructural, with Ni (Cu) and O ions arranged in square planar coordination within NiO_2 (CuO_2) planes that are stacked along the c axis. In case of cuprates, even without IL structure, these planes crucially host the superconducting condensate, which emerges upon doping with charge carriers when the parent insulating ground state with long-range antiferromagnetic (AFM) order vanishes.

However, it was realized early that the effective electronic structures of IL nickelates and cuprates are distinct to a certain degree [27]. Notably, x-ray spectroscopic measurements demonstrated that $3d$ - $2p$ hybridization between Ni and its O ligands is minimal in IL $(\text{La}, \text{Nd})\text{NiO}_2$ [28,29], while strong $3d$ - $2p$ mixing of Cu and O is a hallmark of cuprates. Instead, the presence of states with mixed Ni $3d$ and rare-earth $5d$ character was detected [28,29], which indicates a distinguished role of the rare-earth spacer layer and was also predicted in *ab initio* calculations [18,27,30–33]. While it is still under debate whether the hybridized three-dimensional states of weakly interacting $5d$ electrons play an active role in shaping the low-energy physics of IL nickelates [19,32,34–38], recent spectroscopic studies reported that the doped holes in $\text{Nd}_{1-x}\text{Sr}_x\text{NiO}_2$ reside predominantly in the planar Ni $3d$

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orbitals [39], giving rise to a $3d^8$ spin-singlet state of hole pairs [40].

Yet, a coherent picture of the magnetism in IL nickelates has not been established. In fact, previous theoretical approaches often addressed magnetic correlations in terms of static AFM states [26,27,31,35,41–45], while such long-range ordered states have remained elusive in experiments [5,22,23,46–51]. This absence is in stark contrast to superconducting cuprates, where the undoped (parent) compounds exhibit commensurate AFM order [2]. Notably, strong AFM correlations persist in cuprates in the form of spin fluctuations [52] even when charge-carrier doping suppresses long-range order [2]. Prominently, the signatures of such fluctuations were detected in doped cuprates in the form of paramagnons [53], which are damped spin excitations that pervade the phase diagram up to highest doping levels [54,55]. Conversely, it was revealed that similar damped spin excitations emerge in undoped IL nickelates [23]. Thus, it is a pressing issue to clarify the nature of magnetic correlations in IL nickelates with respect to the phenomenology of cuprates in a concerted effort between experiment and theory.

In addition to the comparison to other unconventional superconductors, IL nickelates are a material class that is intriguing in its own right. For instance, it was recently proposed [19] that IL nickelates could be a genuine realization of the single-band Hubbard model [56–61], which is arguably the most paradigmatic model for correlated electron systems. This would allow one to describe long- and short-ranged spatial correlations by means of cutting-edge quantum-field theoretical methods. Along these lines the dynamical vertex approximation (D Γ A) [62,63], which is a diagrammatic extension [64] of the dynamical mean-field theory (DMFT) [65–67], was recently applied to a single-band low-energy representation of Nd-based IL nickelates [19]. The D Γ A calculations yielded good agreement with the experimentally observed T_c of 10–15 K in Nd_{0.8}Sr_{0.2}NiO₂ thin films [5] which motivates our explicit analysis of the magnetic correlation functions in the same theoretical framework. Additionally, the setting of $s = 1/2$ spins on a square lattice was predicted to bear extraordinary spin correlations [68–70] even in absence of long-range order [71–73]. Specifically for IL nickelates, proximity to a frustrated quantum critical point was proposed [74] that could give rise to a conducting spin-liquid phase [16,74], while other theoretical studies have suggested the occurrence of a spin-freezing crossover due to pronounced Hund's coupling [13].

In this work, we shed light on the magnetic correlations in polycrystalline LaNiO₂ using specific heat (C_p), muon-spin rotation (μ SR), and static (χ) and dynamic (χ' , χ'') magnetic susceptibility measurements. Along the lines of previous experiments [46–48], we obtained the IL phase from topotactic reduction of LaNiO₃ in the perovskite phase. In addition, LaNiO₃ was employed as a reference for the analysis of C_p and the susceptibilities of LaNiO₂. In the C_p of LaNiO₂ we observe residual entropy at low temperatures, indicating the presence of spin fluctuations and related paramagnon excitations. Our μ SR and low-field static and dynamic magnetic susceptibility measurements reveal short-range magnetic correlations and cluster glasslike spin freezing

at low temperatures. The glassy dynamics are attributed to subtle nonstoichiometries of the oxygen sublattice, which develop during the synthesis of LaNiO₂ via topochemical reduction of LaNiO₃. The corrected intrinsic susceptibility χ_{corr} of LaNiO₂ can be extracted from susceptibility measurements in strong external fields, unveiling a distinct non-Curie-Weiss behavior over a wide temperature range. In particular, we find that $\chi_{\text{corr}}(T)$ continuously increases with increasing temperature, which is reminiscent of the susceptibility of underdoped cuprates. This temperature dependence of χ_{corr} can be qualitatively understood by a multimethod study of the two-dimensional single-band Hubbard model using two complementary quantum field-theoretical methods, dynamical mean-field theory (DMFT), cellular dynamical mean-field theory (CDMFT), and the dynamical vertex approximation (D Γ A). In the context of this theoretical understanding of the nature of magnetic correlations in IL nickelates, we also comment on the connection between the obtained non-Curie-Weiss behavior and the opening of a pseudogap.

II. METHODS

A. Experimental methods

Polycrystalline LaNiO₃ powder was synthesized via the citrate-nitrate method [75]. The metal nitrates Ni(NO₃)₂ and La(NO₃)₃ and citric acid were dissolved in water. The homogeneous solution was heated on a hotplate to 500 °C, which evaporates the water and finally decomposes the nitrates. The oxide was further treated by repeated dry ball-milling and calcination at 750 °C for at least 12 h. Subsequently, the powder was treated in an autoclave with 400-bar O₂ pressure to ensure phase purity and full oxygenation. The autoclave starting temperature of 450 °C was decreased to 250 °C over 2 days. The LaNiO₂ IL phase was obtained through topotactic reduction of the LaNiO₃ perovskite phase using CaH₂ as a reducing agent. Batches of 50 mg LaNiO₃ powder were wrapped in aluminum foil with an opening at one end and loaded into quartz tubes with approximately 250 mg CaH₂ powder. To prevent the CaH₂ from reacting with air, the procedure was performed in an Ar-filled glove box and the quartz tubes were sealed under vacuum ($\approx 10^{-7}$ mbar). After sealing, the tubes were heated in a furnace with a ramp rate of 10 °C/min and kept at 280 °C for 316 h. Finally, the samples were cooled at a similar rate and the reduced powder was then extracted. During the entire process, the nickelate and CaH₂ powder have not been in direct contact.

LaNiO₃ and the reduced phase were characterized by powder x-ray diffraction (PXRD) (see Appendix A). The refined structural parameters (Table I) are in good agreement with previous reports [47,48,76]. Furthermore, the refinement of LaNiO₃ [Fig. 7(a)] indicates that the citrate-nitrate method and subsequent autoclave treatment yields highly pure powders without traceable secondary phases. Also the data on the reduced sample can be refined assuming a single phase of IL LaNiO₂ after a reduction time of 316 h [Fig. 7(b)], without indications for impurities due to incomplete reduction or decomposition, such as LaNiO_{2.5}, NiO/Ni, or La₂O₃. For shorter reduction times, residues of phases with excess oxygen were detected, for instance, we found that after

196 h the reduced sample contained 14.7 wt.% of $\text{LaNiO}_{2.5}$. The phase purity was further confirmed on pressed LaNiO_3 and LaNiO_2 pellets by scanning electron microscopy with energy dispersive x-ray spectroscopy (EDS) and no agglomerations of elemental Ni were found (see Appendix A). Inductively coupled plasma mass spectrometry (ICP-MS) indicated ideal stoichiometries (within the experimental error) of $\text{La}_{0.99(1)}\text{Ni}_{0.99(1)}\text{O}_{2.99(3)}$ and $\text{La}_{0.99(1)}\text{Ni}_{0.99(1)}\text{O}_{2.02(3)}$, respectively (see Appendix A).

The specific-heat data were collected with the standard options of a Physical Property Measurement System (PPMS, Quantum Design) using a thermal relaxation method on cold-pressed pellets of LaNiO_3 and LaNiO_2 powder. The low-temperature dc susceptibility measurements were performed using a vibrating sample magnetometer (MPMS VSM SQUID, Quantum Design) and the ac susceptibility measurements using a Physical Property Measurement System (PPMS, Quantum Design) with an ACMS option. High-temperature magnetization measurements were carried out in a Quantum Design MPMS3 setup.

The μSR experiments were performed on the General Purpose Surface-Muon Instrument (GPS) [77] at the Paul Scherrer Institute (PSI) in Switzerland. The GPS instrument was operated in veto mode, which minimizes the background signal below the detection limit. The data analysis was performed with the free software package MUSRFIT [78].

B. Theoretical methods

For the theoretical modeling of LaNiO_2 , we employ the parameters given in a recent single-band Hubbard model study for NdNiO_2 [19]: We set the onsite Coulomb interaction to $U = 3.2$ eV, and the tight-binding parameters to $t = 395$ meV, $t' = -95$ meV, and $t'' = 47$ meV (expressed via the nearest-neighbor transition amplitude t , these parameters read as $U = 8t$, $t' = -0.25t$, and $t'' = 0.12t$). Like for NdNiO_2 , we neglect a (small) hopping integral in c direction also for LaNiO_2 . Note that the electronic structure of LaNiO_2 corresponds more closely to a half-filled $d_{x^2-y^2}$ band than NdNiO_2 [19], suggesting that the half-filled model is even more appropriate for the former case. Hence, we use a half-filled model for all calculations shown.

In order to obtain the magnetic susceptibility we apply multiple many-body techniques for correlated systems to the same model. Such multimethod studies have proven highly valuable [79–81] for the calculation of observables in challenging parameter regimes of the model. Here we employ the dynamical mean-field theory (DMFT [65–67]) and two complementary extensions thereof: (i) the dynamical vertex approximation (D Γ A [62,63]), which is a diagrammatic extension [64] of DMFT, and the cellular dynamical-mean field theory (CDMFT), a real-space cluster extension [82] of DMFT. We apply the D Γ A in its ladder version with Moriyaesque λ corrections in the spin channel [83] and the CDMFT on a $N_c = 4 \times 4 = 16$ site cluster [82,84].

DMFT includes all local quantum fluctuations of the system, however, entirely neglects spatial correlations. CDMFT captures nonlocal correlations exactly up to the cluster size. Cluster extensions of DMFT exhibit a non-Curie-Weiss behavior in the susceptibility [85–87], associated with the onset

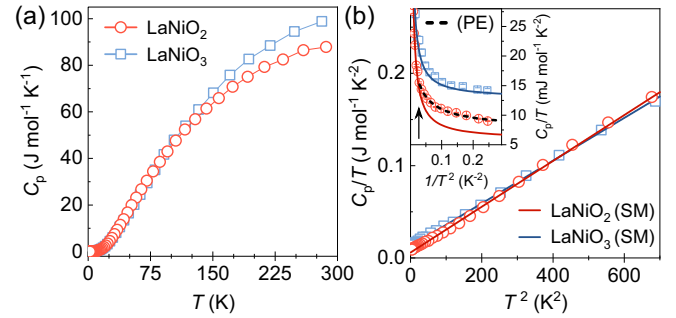


FIG. 1. (a) Specific heat C_p of polycrystalline LaNiO_2 (red) and LaNiO_3 (blue). (b) C_p/T as a function of T^2 . Solid lines are standard model (SM) fits of LaNiO_2 and LaNiO_3 between $T = 6$ and 25 K according to $C_p = \gamma T + \beta T^3$ (see text). The inset shows C_p/T as a function of $1/T^2$. The black arrow centered at $T = 6$ K indicates the onset of a deviation from the extrapolated SM fit (solid red line) for LaNiO_2 . The dashed black line is a model fit of LaNiO_2 between $T = 2$ and 6 K according to $C_p = \gamma T + \beta T^3 + \alpha T^3 \ln(T)$, which accounts for an extra contribution due to paramagnon excitations (PE).

of the pseudogap regime of cuprates [88,89]. These methods are controlled and unbiased with respect to fluctuation channels. Let us note that a non-Curie-Weiss behavior was also recently observed in a minimally entangled typical thermal states study on finite cylinders [90] as well as in a diagrammatic Monte Carlo study in the weak-coupling regime of the Hubbard model [91].

Our complementary method, the D Γ A, is able to include short- and long-range spatial correlations of the chosen (magnetic) channel on equal footing. The D Γ A has proven particularly successful in the vicinity of second-order phase transitions [92–94], for the estimation of superconducting transition temperatures [19,95,96], for obtaining insights into the physics of layered materials and dimensional crossovers [97,98], and in comparison to numerically exact benchmark calculations in the weak-coupling regime of the Hubbard model [80]. The (isolated) finite-size systems have been diagonalized exactly with PYED in the TRIQS [99] package. For further calculational details, see Appendix E.

III. RESULTS

A. Specific heat

Nickelates in the IL phase can be obtained via the topotactic removal of oxygen from a precursor $R\text{NiO}_3$ ($R = \text{La}, \text{Pr}, \text{or Nd}$) perovskite phase [5,46–51]. In this work, we reduce highly pure LaNiO_3 powder to LaNiO_2 (see Methods). Possible admixture of secondary phases lies below the detection threshold of our PXRD characterization (see Appendix A).

As a first step to explore magnetic correlations in LaNiO_2 we determine the temperature dependence of the specific heat (C_p) between 2 and 297 K (Fig. 1). Characteristic anomalies in C_p not only reveal the presence of phase transitions, but can also provide valuable information in absence of long-range order, for instance about spin fluctuations [100–102] or frustrated spin states [103]. As expected, λ -like anomalies that correspond to magnetic and/or structural phase transitions are

absent in $C_p(T)$ of LaNiO_2 [Fig. 1(a)]. This is consistent with neutron diffraction experiments on polycrystalline LaNiO_2 and NdNiO_2 that reported a lack of magnetic Bragg reflections [48,49]. A qualitative understanding of C_p of LaNiO_2 can be gained from a direct comparison to the precursor phase LaNiO_3 , whose C_p is remarkably similar at first glance [Fig. 1(a)], in spite of different crystallographic symmetries (Table I). In particular, below 120 K the specific heat of the two compounds is almost indistinguishable, which is likely related to the similarity of their La-Ni sublattice (Table I), which gives the major contribution to C_p in this temperature range. In contrast, phonons involving the oxygen sublattice, which is substantially altered upon reduction, mostly play a role at higher temperatures and can be responsible for the slightly larger C_p of LaNiO_3 above 120 K [Fig. 1(a)].

Figure 1(b) shows the specific heat below 25 K, which in case of LaNiO_3 can be described by $C_p = \gamma T + \beta T^3$ [104], according to the standard model for a nonmagnetic solid. A fit between 6 and 25 K results in a Sommerfeld coefficient $\gamma = 12.2(4) \text{ mJ mol}^{-1} \text{ K}^{-2}$, a lattice contribution $\beta = 0.234(8) \text{ mJ mol}^{-1} \text{ K}^{-4}$, and a Debye temperature of $\Theta_D = 346 \text{ K}$, in good agreement with previous studies on polycrystalline LaNiO_3 [104]. For LaNiO_2 , the same fit yields $\gamma = 4.4(6) \text{ mJ mol}^{-1} \text{ K}^{-2}$, $\beta = 0.255(6) \text{ mJ mol}^{-1} \text{ K}^{-4}$, and $\Theta_D = 336 \text{ K}$. Thus, the lattice contribution β remains almost unchanged in comparison to LaNiO_3 , whereas the electronic contribution γ of LaNiO_2 decreases significantly. In general, such decrease can be related to a less correlated electronic character and/or reduced metallicity, which is still an open question for LaNiO_2 : Electrical transport measurements on high-quality epitaxial LaNiO_2 thin films revealed a metallic in-plane conductivity [105], whereas powder measurements possibly suffer from extrinsic effects, such as inferior contacts between grain boundaries, and the anisotropy between in-plane and c -axis transport, rationalizing the observed non-metallic behavior [48].

The inset in Fig. 1(b) focuses on the evolution of the specific heat towards the lowest measured temperatures and shows C_p/T as a function of $1/T^2$. As expected for the nonmagnetic Fermi-liquid LaNiO_3 [106], we find that the data below 6 K are well described by an extrapolation of the standard model fit $C_p = \gamma T + \beta T^3$ performed between 25 and 6 K. However, a similar extrapolation of a standard model fit of LaNiO_2 does not capture the measured data below 6 K [inset in Fig. 1(b)]. This deviation is small in absolute terms, but nevertheless statistically significant. While we cannot exclude that this residual entropy emerges due to disorder in the polycrystalline powder, we emphasize that such an additional contribution to C_p can typically arise from the emission and reabsorption of persistent spin fluctuations or damped spin waves (paramagnons) [100–102]. Employing a large- U Hubbard model with parameter choices corresponding to the paramagnetic, metallic regime at half-filling, the spin fluctuation term for the specific heat follows as $T^3 \ln(T)$ [102]. Thus, we fit the specific heat of LaNiO_2 between 2 and 6 K by $C_p = \gamma T + \beta T^3 + \alpha T^3 \ln(T)$ and obtain $\gamma = 7.7(9) \text{ mJ mol}^{-1} \text{ K}^{-2}$, $\beta = 0.5(1) \text{ mJ mol}^{-1} \text{ K}^{-4}$, and $\alpha = -0.16(1) \text{ mJ mol}^{-1} \text{ K}^{-4}$. Since such fit provides an adequate description of the data below 6 K, it can be an indication of the presence of paramagnons in LaNiO_2 . Notably, paramagnons

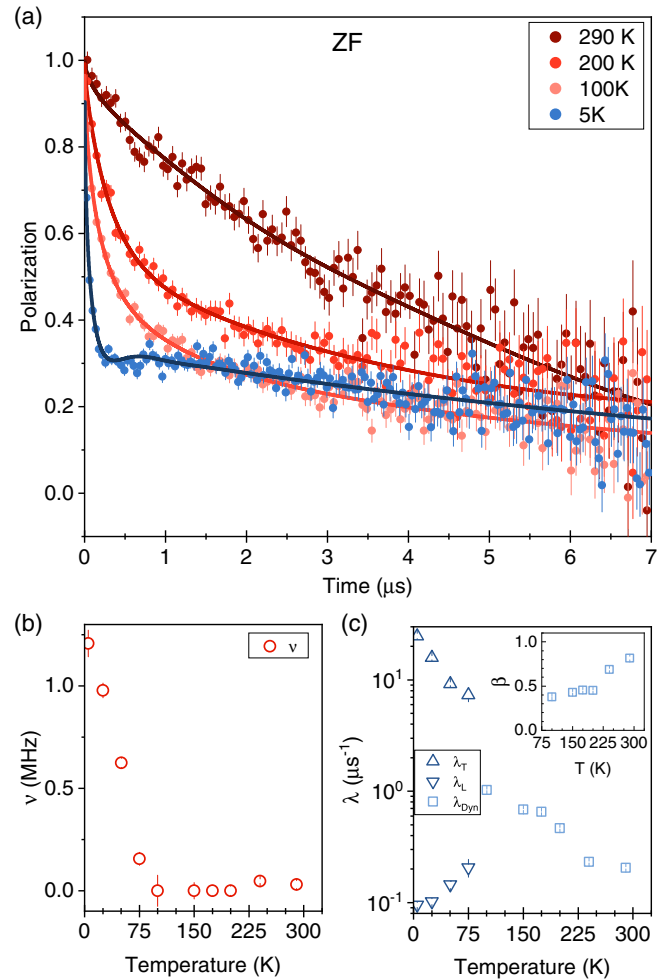


FIG. 2. (a) Zero-field (ZF) μSR spectra of LaNiO_2 at representative temperatures. Solid lines are fits of the polarization $P(t)$ (see text). (b) Temperature dependence of the parameter ν , which is indicative of the internal magnetic field. (c) Temperature dependence of the parameters λ_L , λ_T , and λ_{Dyn} , extracted from fits to regimes below 75 K and above 100 K, respectively. The inset shows the evolution of the parameter β for temperatures above 100 K obtained from a simple stretched exponential fit.

are not only a hallmark of doped cuprates [53,54], but damped spin excitations in absence of long-range order were recently also detected with resonant inelastic x-ray scattering (RIXS) in thin films of the IL nickelate NdNiO_2 [23] and the structurally related trilayer nickelate $\text{Pr}_4\text{Ni}_3\text{O}_8$ [107].

B. Muon-spin rotation

Insights into quasistatic and dynamical magnetic processes can be gained from μSR , with positive muons stopping on interstitial lattice sites and acting as a sensitive local probe for small internal magnetic fields and ordered magnetic volume fractions in a sample. Figure 2(a) displays representative zero-field (ZF) μSR spectra of LaNiO_2 for temperatures between 290 and 5 K. The full set of ZF μSR spectra are shown in Fig. 9. In accordance with the absence of a magnetic transition in our C_p measurements, the μSR spectra do not show well-defined oscillations that would be

indicative of long-range magnetic order, as observed for instance in the AFM trilayer nickelate $\text{La}_4\text{Ni}_3\text{O}_8$ [108]. Nonetheless, complementary measurements in longitudinal fields (LF) at 295 K reveal that fluctuating moments are present in the sample (Fig. 10). The ZF spectra below 75 K [Figs. 2(a) and 9] exhibit a strongly damped oscillation with a first local minimum around $0.25 \mu\text{s}$, suggesting the occurrence of pronounced short-range order at these temperatures. Moreover, the long-time tail of the 5-K spectrum lies above the 100-K spectrum and approaches a value of $\frac{1}{3}$ of the initial polarization, which signals quasistatic magnetism as observed, for instance, below the spin-freezing transition in spin glasses [109]. Figure 2(b) shows the ZF μSR frequency $\nu(T)$ from fitting the ZF data as described below, suggesting a crossover with an onset between 75 and 100 K, which we attribute to the onset of glassy behavior, where magnetic moments freeze at low temperatures with random orientations while long-range ferromagnetic or AFM order remain absent [110]. We consider two fitting regimes for the data, i.e., purely dynamic behavior $P(t) = P_0 e^{-(\lambda_{\text{dyn}} t)^\beta}$ above 75 K, and slowly fluctuating and quasistatic behavior which is described by $P(t) = 1/3 e^{-\lambda_L t} + 2/3 \cos(2\pi \nu t) e^{-(\lambda_T t)^\beta}$ [Fig. 2(c)] below this temperature. Here, λ_T and λ_L are the transverse and longitudinal relaxation rates. The $\frac{2}{3}$ (transverse) and $\frac{1}{3}$ (longitudinal) components reflect the polycrystalline nature of the sample leading to a powder average of the internal fields with respect to the initial muon spin direction.

Figure 2(c) and the inset in Fig. 2(c) show the temperature evolution of the muon spin relaxation rate λ and the stretch parameter β in the dynamic regime between 295 and 75 K, respectively. The observed increase of λ with decreasing temperature and variation of β from approximately 1 to 0.5 indicating a broad distribution of magnetic relaxation times typical for a spin glass prior to the freezing transition

[109,111,112]. Notably, the temperature range across which this variation occurs is relatively broad. This is a typical feature of heterogeneous samples, with the muons experiencing both strong and small fields in the sample. Thus, polycrystalline LaNiO_2 likely exhibits a spatially inhomogeneous distribution of magnetic correlations, possibly due to the presence of magnetic clusters or islands. Fluctuations of spatially separated correlated clusters are typically slow, which is compatible with the extracted timescales in Fig. 2(c). Below 75 K the dynamic relaxation rate λ_L decreases consistent with a continuous slowing down also in the quasistatic regime. The observed peak in the dynamic relaxation rate is a fingerprint of magnetic fluctuations slowing down and crossing the time window of the measurement. The transverse relaxation rate λ_T and the frequency ν of the overdamped oscillation increase below 75 K, indicative of the growing magnetic correlations.

In summary, the μSR data suggest the presence of glassy magnetic behavior with an inhomogeneous spatial and temporal distribution. Nevertheless, especially the observation of a ZF μSR oscillation with a large relaxation rate λ_T of up to $25 \mu\text{s}^{-1}$ together with the typical separation of the spectra in a $\frac{1}{3}$ and a $\frac{2}{3}$ component is a strong indication that the magnetic behavior originates from the bulk of the sample.

C. Magnetic susceptibility

As a next step, we investigate the static magnetic susceptibility (χ) of LaNiO_2 . Figure 3(a) shows $\chi(T)$ of LaNiO_2 measured in a relatively small external field of 0.1 T. In accordance with previous experiments [46–49,51] and our C_p and μSR results, the magnetization signal lacks any signatures of long-range magnetic ordering between 4 and 300 K. However, we observe several salient features, which were not reported in early studies of polycrystalline LaNiO_2

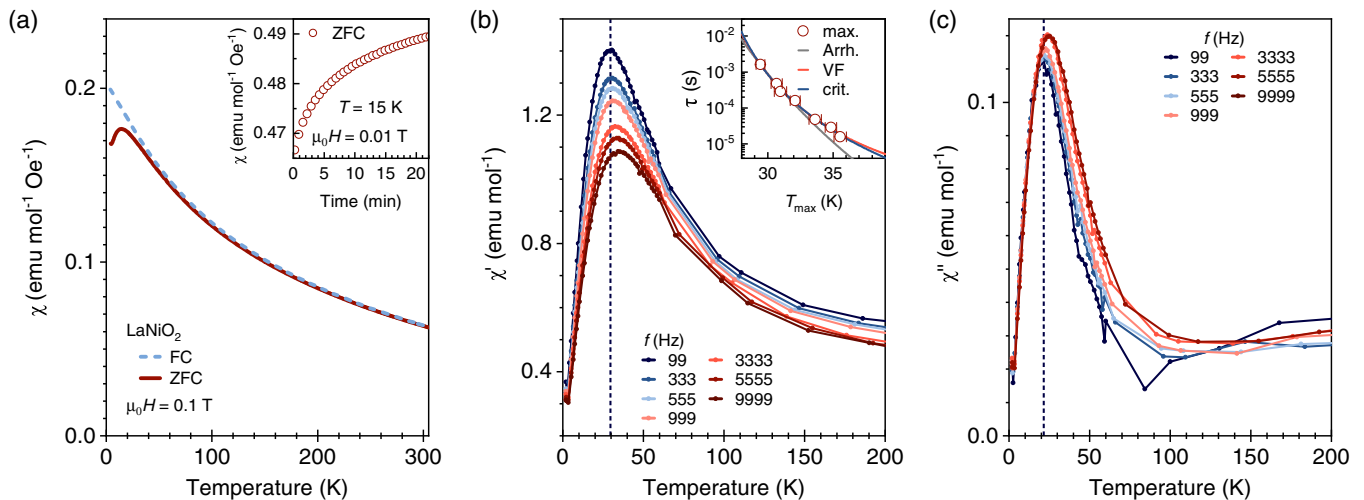


FIG. 3. Magnetic susceptibility of LaNiO_2 in small external fields. (a) Static susceptibility χ measured upon heating after zero-field cooling (ZFC, solid line) and field-cooling (FC, dashed line), respectively, in an external field of $\mu_0 H = 0.1$ T. The inset shows the evolution of χ (ZFC, $\mu_0 H = 0.01$ T) as a function of time for an intermittent stop at $T = 15$ K. (b) Real part of the dynamic susceptibility χ' for different ac drive frequencies f . The static external field is zero and the maximum field amplitude of the ac field is 4×10^{-4} T. Vertical dashed lines indicate the positions of the maximum for $f = 99$ Hz. The inset shows the relation between the temperature T of the maximum of χ' and the time period $\tau = (2\pi f)^{-1}$ of the ac field. The solid gray, red, and blue lines are fits with an Arrhenius (Arrh.), a Vogel-Fulcher (VF), and a critical dynamical scaling (crit.) law, respectively. (c) Imaginary part of the dynamic susceptibility χ'' .

[46,48]. In particular, we detect an irreversibility between the zero-field-cooled (ZFC) and field-cooled (FC) magnetization branches at low temperatures and a cusplike feature of the ZFC branch centered around 17 K. Such behavior of the ZFC and FC magnetization is characteristic for spin glasses. Furthermore, hysteretic behavior is observed in measurements of the isothermal magnetization [Fig. 11(b)], which is enhanced at temperatures corresponding to the region of the cusp of the ZFC curve and below. Closely similar ZFC-FC splittings and magnetic hysteresis were also reported in recent studies on polycrystalline $R\text{NiO}_2$ and $\text{Pr}_4\text{Ni}_3\text{O}_8$ powders [113,114], as well as $\text{La}_{1-x}\text{Ca}_x\text{NiO}_{2+\delta}$ crystals [115]. In principle, the maximum of the cusp can be associated with the freezing temperature T_f of the glass; however, note that our measurement of $\chi(T)$ does not probe the equilibrium state of the spins. Specifically, the cusp feature of the ZFC curve shifts to higher temperatures when the $\chi(T)$ measurements are conducted in smaller external static fields [Fig. 11(a)]. Moreover, χ evolves as a function of the measurement time [inset in Fig. 3(a)], which is an additional indication that dynamical processes are at play. Thus, the observed maximum of the cusp at 17 K in Fig. 3(a) is only an approximate indicator of T_f .

Detailed insights into the dynamics and the nature of the spin-glass phase can be gained from measurements of the real and imaginary parts of the susceptibility χ' and χ'' in an ac drive field. Figures 3(b) and 3(c) show that the ac susceptibilities of LaNiO_2 exhibit a pronounced cusp/peak at low temperatures, which is reminiscent of the cusp in the dc susceptibility [Figs. 3(a) and 11(a)]. We note that in particular the onset of the peak in $\chi''(T)$ [Fig. 3(c)] coincides with the onset of the spin freezing identified in the μSR measurements (Fig. 2). With increasing drive frequency f , the maximum of the peak in $\chi'(T)$ shifts to higher temperatures and its amplitude decreases. Similarly, the position of the peak in $\chi''(T)$ increases, whereas its amplitude and width also increase with increasing f [Fig. 3(c)]. Such behavior of χ' and χ'' is typical for both canonical and cluster spin glasses [110], while the μSR results strongly indicated spatial inhomogeneities, i.e., our sample consists of an assembly of interacting magnetic clusters.

The interaction strength between magnetic clusters can be assessed from the scaling behavior [110] of the frequency-dependent peak position of $\chi'(T)$ [Fig. 3(b)]. To this end, the frequency-dependent peak position in χ' is fitted using different laws [inset in Fig. 3(b)], which are appropriate for increasing interaction strengths, including an Arrhenius, a Vogel-Fulcher (VF), and a critical slowing-down law. An Arrhenius law can capture systems of low-dimensional noninteracting magnetic entities and is expressed as $\tau = \tau_0 \exp(E_a/k_B T)$, where $\tau = (2\pi f)^{-1}$ is the relaxation time, τ_0 a characteristic time, E_a an energy barrier separating two low-energy states, and k_B the Boltzmann constant. Here, we obtain $\tau_0 = 4.6 \times 10^{-17}$ s and $E_a/k_B = 918$ K. The nonphysically small value of τ_0 and the extremely large E_a [110] suggest that LaNiO_2 does not fall into the noninteracting Arrhenius limit. In addition, we can rule out superparamagnetic behavior, which exhibits typical values of τ_0 between 10^{-9} and 10^{-13} s [110,116]. In presence of moderate intercluster couplings in a spin glass, a VF law according to $\tau = \tau_0 \exp[E_a/k_B(T - T_0)]$ can be employed, where T_0 is the glass temperature, which

typically is similar to the freezing temperature. For LaNiO_2 , we obtain $\tau_0 = 1.2 \times 10^{-8}$ s, $E_a/k_B = 118.4$ K, and $T_0 = 19.4$ K. This yields $E_a/(k_B \tau_0) \approx 6$, indicating that LaNiO_2 is located in the weak to intermediate coupling regime, whereas $E_a/(k_B \tau_0) \ll 1$ would correspond to strong coupling. The obtained timescale τ_0 is comparable with the relaxation rate λ_L extracted from the μSR measurements, suggesting that the same freezing phenomenon is probed by the two complementary techniques. For completeness, we also test a strongly interacting critical dynamical scaling law according to a power law of the form $\tau = \tau_0[(T - T_f)/T_f]^{-z\nu}$, with the freezing temperature T_f in the limit $f \rightarrow 0$ and the dynamic critical exponent $z\nu$ that accounts for a critical slowing down of the dynamics of the magnetic entities in proximity to T_f . A fit of the data in the inset in Fig. 3(b) results in $\tau_0 = 1.6 \times 10^{-6}$ s, $T_f = 19.5$ K, and $z\nu = 10.8$. While the critical exponent adopts an acceptable value, τ_0 is larger than that of typical canonical and cluster glasses [110]. In consequence, our polycrystalline LaNiO_2 sample is best described by a VF law with weak to intermediate coupling between magnetic clusters. We note that a VF law was also employed to fit the susceptibility of polycrystalline $\text{Pr}_4\text{Ni}_3\text{O}_8$ [114] and $R\text{NiO}_2$ [113], although a stronger coupling $E_a/(k_B \tau_0)$ was reported.

The observation of glassy behavior with weak to intermediate coupling strength in the μSR and dynamic susceptibility measurements raises the question whether glassiness is an intrinsic property of LaNiO_2 , or arises from possible impurities or secondary phases. In the following, we address this issue by susceptibility measurements in strong external fields (Fig. 4). In the case of ferromagnetic impurities in reduced powder samples [117,118], it is expected that their magnetization signal saturates in sufficiently strong fields, yielding only a constant contribution after the field exceeds a certain threshold value [48]. Moreover, large fields suppress the glassy dynamics [Fig. 11(a)] and underlying intrinsic magnetic correlations can be exposed. Figure 4(a) displays the static susceptibility χ of our polycrystalline LaNiO_2 sample in a field of 7 T. In contrast to the low-field measurement [Fig. 3(a)], an irreversibility between the ZFC and FC curves is not present, thus confirming that spin-glass effects do not dominate the signal at this field strength. Furthermore, χ does not depend on measurement time or history [inset of Fig. 4(a)]. Overall, χ increases almost linearly with decreasing temperature, while a subtle upturn at lowest temperatures and a broad hump centered around 200 K can be identified. To determine whether these features are related to the intrinsic paramagnetic susceptibility of LaNiO_2 , or induced by (ferromagnetic) impurities, we measured magnetization-field isotherms $M(H)$ at various temperatures [Fig. 4(b)]. The isotherms show a pronounced nonlinearity for small applied fields, which decreases for isotherms measured at higher temperatures and vanishes between 600 and 650 K. Since the Curie temperature of Ni is ~ 620 K, the nonlinearity is likely due to the presence of a small amount of Ni impurities, which we estimate to be less than 2.5 wt.% (see Appendixes A and C). The saturation magnetization M_{sat} can be determined from linear fits to the isotherms at high fields in the range between 5 and 7 T (see Appendix C). The temperature dependence of M_{sat} [Fig. 4(c)] qualitatively resembles $\chi(T)$ in Fig. 4(a), suggesting the temperature dependence of $\chi(T)$ in Fig. 4(a) is dominated by the

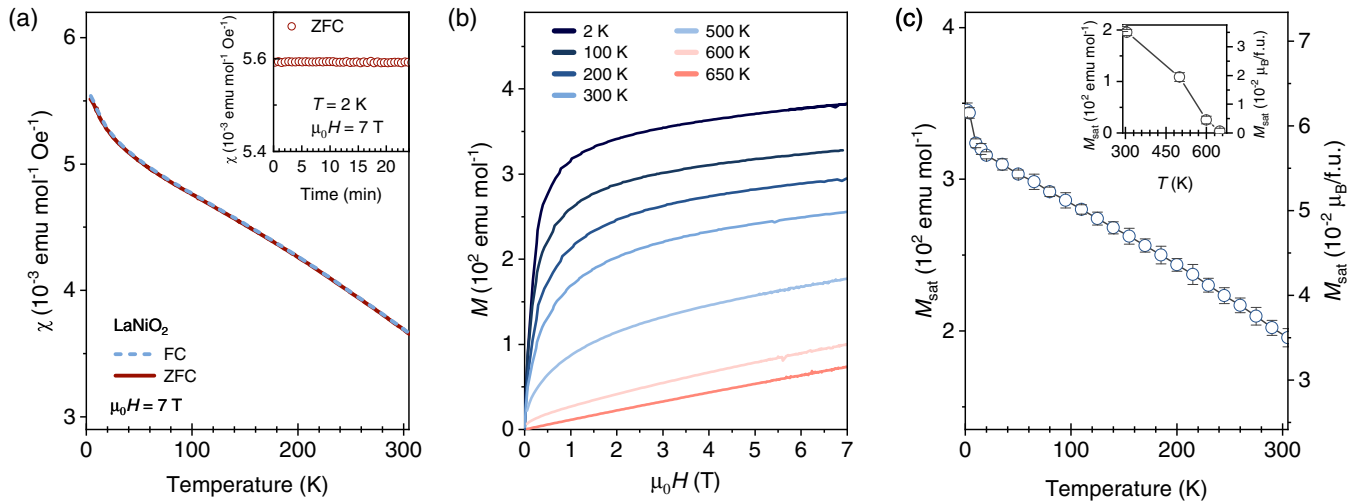


FIG. 4. Magnetic susceptibility of LaNiO_2 in strong external fields. (a) ZFC and FC susceptibility χ in an external field $\mu_0 H = 7$ T. The inset shows the evolution of χ (ZFC) as a function of time for an intermittent stop at $T = 2$ K. (b) Isothermal magnetization between 0 and 7 T at representative temperatures. (c) Saturation magnetization M_{sat} extracted from linear fits of the isothermal magnetization curves between 5 and 7 T for temperatures between 2 and 305 K (see Appendix C). The inset shows M_{sat} extracted from isotherms measured at higher temperatures.

signal of the ferromagnetic Ni impurities and does not reflect the intrinsic susceptibility of LaNiO_2 .

Nevertheless, a distinction between the ferromagnetic Ni background and intrinsic paramagnetic behavior of the majority phase is possible by applying the Honda-Owen method [119,120] to the isotherms measured between 5 and 7 T (see Appendix C). More specifically, the Honda-Owen method extrapolates the measured susceptibility $M/H = \chi_{\text{corr}} + C_{\text{sat}} M_{\text{sat}}/H$ for $1/H \rightarrow 0$, where M/H is the measured susceptibility, χ_{corr} the corrected, intrinsic susceptibility, C_{sat} the presumed ferromagnetic impurity content, and M_{sat} its saturation magnetization (see Appendix C). Figure 5 shows the temperature dependence of χ_{corr} , which is drastically different from $\chi(T)$ in Fig. 4(a). Specifically, $\chi_{\text{corr}}(T)$ exhibits a minimum around 65 K and increases almost linearly towards room temperature, signaling a strik-

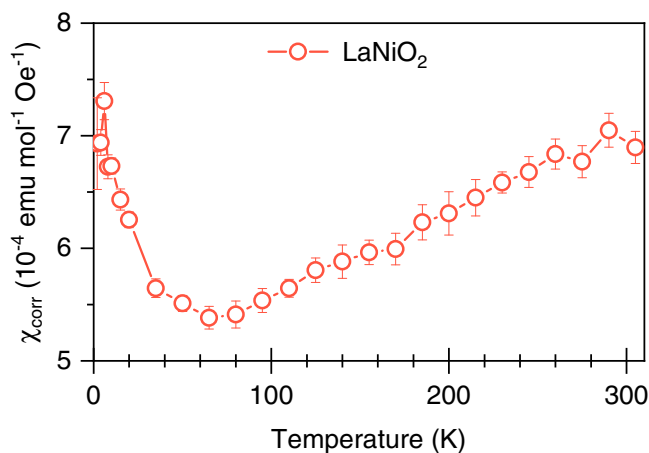


FIG. 5. Corrected paramagnetic susceptibility χ_{corr} extracted from isothermal magnetization curves by the Honda-Owen method (see Appendix C).

ing non-Curie-Weiss behavior. We note that this increase is qualitatively consistent with the temperature dependence of the magnetic Knight shift in a recent ^{139}La NMR study on polycrystalline LaNiO_2 [22], confirming that $\chi_{\text{corr}}(T)$ is not dominated by a spurious signal of Ni-related impurities. The monotonous increase of $\chi_{\text{corr}}(T)$ continues at least up to 600 K (Fig. 13). Remarkably, such behavior of $\chi_{\text{corr}}(T)$ closely resembles the susceptibility of doped cuprates, such as $\text{La}_{2-x}\text{Sr}_x\text{CuO}_4$ with $x \geq 0.04$ [121]. Furthermore, a monotonous increase of the susceptibility was also reported for iron pnictide superconductors and attributed to local AFM correlations [122]. In contrast to this monotonously increasing susceptibility, previous studies on polycrystalline LaNiO_2 [48] reported paramagnetic Curie-Weiss behavior between 6 and 300 K, including a crossover between two Curie-Weiss regimes at 150 K. This discrepancy could be due to different amounts of impurity phases in the sample of Ref. [48]. In particular, we determine a shorter c -axis lattice parameter for our LaNiO_2 phase (Table I), suggesting the presence of less excess apical oxygen, possibly due to our longer reduction times (see Methods) and/or the use of CaH_2 instead of NaH as a reducing agent. Nevertheless, our extracted absolute value of χ_{corr} at 305 K of $\approx 6.89 \times 10^{-4} \text{ emu Oe}^{-1} \text{ mol}^{-1}$ is comparable to that of Ref. [48], and also similar to the susceptibility of perovskite LaNiO_3 [123]. Furthermore, $\chi_{\text{corr}}(T)$ exhibits an upturn below 65 K (Fig. 5), which was similarly reported in Ref. [48] and is likely related to paramagnetic impurities.

D. Theoretical modeling

The observed non-Curie-Weiss behavior hints towards the relevance of local and nonlocal spin fluctuations as a possible origin of the unusual magnetic response of LaNiO_2 . In the following, we therefore go beyond effective single-Slater-determinant theories, such as the DFT+ U approaches of previous studies [26–28,31,35,41–43,45], and turn to quantum

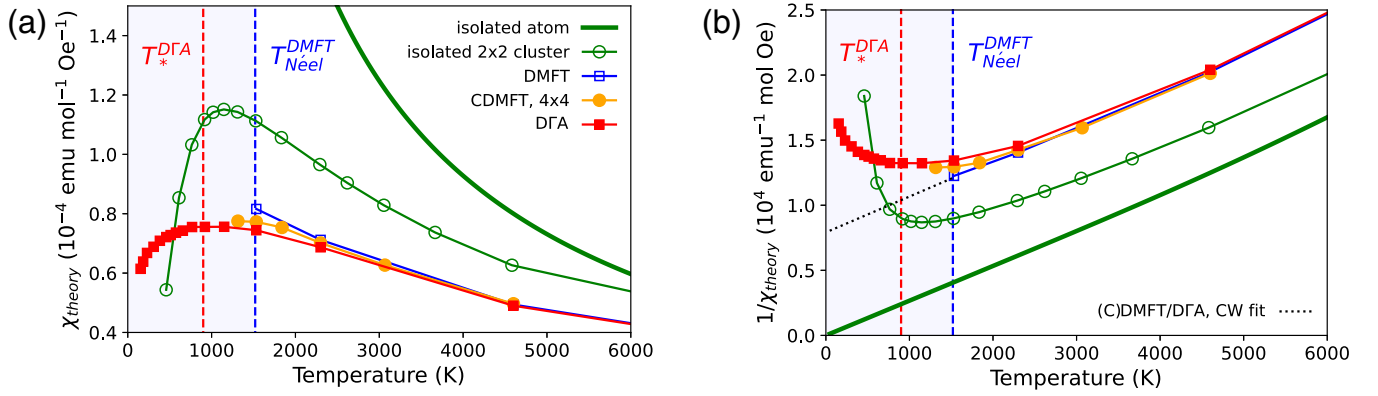


FIG. 6. (a) dc susceptibility χ_{theory} and (b) its inverse calculated for a two-dimensional single-band Hubbard model of LaNiO_2 by various many-body techniques. CDMFT (filled orange circles) and D Γ A (filled red squares) exhibit a deviation from Curie-Weiss behavior at low T when significant nonlocal correlations set in [the dotted black line in (b) shows a respective Curie-Weiss fit from the high- T regime of all three techniques]. This regime is already signaled in DMFT as an (antiferromagnetically) ordered phase for $T < T_{\text{Néel}}^{\text{DMFT}}$ (blue shaded region). For comparison to finite-size systems also the atomic values (solid green line, Curie law) and the ones of an isolated 2×2 cluster (open green circles) are shown.

field-theoretical methods tailored at the systematic inclusion of temporal and spatial correlations. From their application to the two-dimensional single-band Hubbard model described in Sec. II B we obtain the uniform dc susceptibility χ_{theory} in zero field (see Appendix E). Figure 6(a) summarizes the results of our calculations as a function of temperature T . Initially, we apply DMFT which exhibits (anti)ferromagnetic ordering at $T_{\text{Néel}}^{\text{DMFT}} \approx 1500$ K as a result of its mean-field nature (see also [44,124] for additional DMFT calculations). Above this temperature the uniform susceptibility χ^{DMFT} (open blue squares) can be accurately described by a Curie-Weiss law as can be inferred from Fig. 6(b), which shows the inverse susceptibility $1/\chi_{\text{theory}}(T)$ and a respective Curie-Weiss fit (dotted black line). The progressive inclusion of spatial correlations on different length scales, however, invalidates this picture: already short-ranged spatial correlations, taken into account by CDMFT on a 4×4 cluster (filled orange circles), lead to a deviation from Curie-Weiss behavior for $T \lesssim T_{\text{Néel}}^{\text{DMFT}}$. These short-ranged spatial correlations also lead to a small reduction of the onset temperature of magnetic ordering with respect to its DMFT value, below which the calculation of χ^{CDMFT} is hindered (see Appendix E for the applied algorithm).

In order to treat correlations larger than the cluster size and eventually enter the low-temperature regime, we turn to D Γ A, which is a diagrammatic extension of DMFT tailored for the systematic inclusion of spatial correlations on all length scales. The susceptibility calculated in D Γ A (filled red squares) qualitatively resembles the experimentally determined susceptibility $\chi_{\text{corr}}(T)$ of LaNiO_2 (Fig. 5) with a broad maximum centered around $T_*^{\text{D}\Gamma\text{A}} \approx 900$ K, emphasizing the onset of a strong deviation from Curie-Weiss behavior below that temperature. Note that the maximum of $\chi^{\text{D}\Gamma\text{A}}$ appears *without* the emergence of long-range magnetic order, which is prohibited at finite temperatures in 2D systems due to the Mermin-Wagner theorem [125,126]. Nevertheless, strong nonlocal magnetic fluctuations (paramagnons) can exist in the paramagnetic phase at finite temperatures as a precursor of a $T_{\text{Néel}} = 0$ transition. These paramagnon contributions are quite naturally included in the D Γ A framework

by its consideration of ladder diagrams in both particle-hole channels [62].

Comparing our calculations with the experimentally determined χ_{corr} of LaNiO_2 presented in Fig. 5, we find a satisfactory qualitative agreement: both, the rising edge of the susceptibility's broad maximum as well as its relative change in amplitude (between the value at the maximum and at 65 K) lie approximately in the same range. This is remarkably accurate given the fact that $\chi^{\text{D}\Gamma\text{A}}$ has been obtained from an effective single-band description. In fact, the same model has been used in the context of superconductivity in NdNiO_2 [19], thus corroborating the applicability of a single-band model to IL nickelate compounds. However, a quantitative comparison of the susceptibilities suggests that overall the experimental response is substantially larger than the calculated one, which will be discussed in more detail below.

IV. DISCUSSION

The spin-glass properties of polycrystalline LaNiO_2 were revealed by μSR and magnetic susceptibility measurements in small external fields. Whether this glassy behavior is intrinsic to IL nickelates, or induced by impurities and/or secondary phases is an intricate issue. In our PXRD characterization, the signal of possible impurities and secondary phases, including elemental Ni and $\text{LaNiO}_{2.5}$, was below the detection threshold. Nevertheless, our magnetization measurements indicated the presence of a small amount of Ni impurities. In principle, Ni or NiO/Ni nanoparticles can show spin-glass behavior [127–129], which could rationalize the observed characteristics in the static and dynamic susceptibility in Fig. 3. However, the observation of a ZF μSR oscillation with a large relaxation rate λ_T of up to $25 \mu\text{s}^{-1}$ together with the typical separation of the spectra into a $\frac{1}{3}$ and a $\frac{2}{3}$ component indicate that the glassy magnetic behavior originates from the bulk of the sample, although with an inhomogeneous spatial distribution. Specifically, the μSR data seem to be incompatible with the scenario that a dilute distribution of Ni nanoparticles induces the observed glassy properties.

Moreover, Refs. [113,114] investigated similar polycrystalline $RNiO_2$ and $Pr_4Ni_3O_8$ powders upon reoxidization and decomposition, concluding that Ni/NiO impurities cannot explain the observed glassy properties.

Nonetheless, it seems unlikely that square-planar nickelates precisely realize the frustrated nearest- and next-nearest-neighbor exchange that is required for an intrinsic square-lattice spin-glass system. Instead, we consider it plausible that local oxygen disorder in form of remaining apical oxygen is responsible for the observed glassy properties. Such excess oxygen can increase the Ni valence towards $2+$, yielding an effective moment and consequently a local magnetic cluster, possibly even polarizing the environment to some extent. Furthermore, also entities with a larger spatial extension could be a candidate for the magnetic clusters, including domain walls between regions with differently oriented NiO_2 planes or minute inclusions of $LaNiO_{2.5}$. While further work is required to decisively clarify the microscopic origin of the glass state in $LaNiO_2$, it is noteworthy that magnetic clusters of varying sizes were also reported in polycrystalline $LaNiO_{2.75\pm\delta}$ [130,131], which could be a precursor of the clusters observed in $LaNiO_2$.

Along these lines, it is insightful to compare the glassy phase of $LaNiO_2$ to the spin-glass regime of cuprates. For instance in $La_{2-x}Sr_xCuO_4$, a glassy phase emerges below a freezing temperature T_f around 5–6 K for hole doping concentrations $0.02 \leq x \leq 0.06$, i.e., between the AFM phase and the onset of the superconducting dome [132–134]. Magnetic resonance experiments attributed the spin-glass phase in cuprates to frozen AFM clusters that arise due to the separation of charges into hole-rich and -poor regions [135]. Thus, the nature of the spin glass in IL nickelates seems to be distinct from the one in cuprates. However, we note that this study does not allow us to fully rule out a charge segregation scenario in IL nickelates and supplemental measurements on $LaNiO_2$ are desirable, for instance with magnetic resonance techniques.

The magnetic susceptibility measured at high temperatures and in strong magnetic fields provided access to the intrinsic magnetic correlations of $LaNiO_2$ (Fig. 5). As a key experimental result we found that the corrected magnetic susceptibility $\chi_{\text{corr}}(T)$ follows non-Curie-Weiss behavior and increases above 65 K. We note that it cannot be fully ruled out that residual contributions from impurities persist in our $\chi_{\text{corr}}(T)$ data, in spite of our analysis via the Honda-Owen method. Hence, while the observed non-Curie-Weiss behavior is a robust feature, the presented quantitative values of χ_{corr} in Fig. 5 should be understood in terms of a general benchmark, whereas samples with less (more) impurities might exhibit smaller (higher) absolute values of χ_{corr} . Importantly, in absence of long-range order [48], a non-Curie-Weiss increase of the susceptibility can be indicative of strong nonlocal spin fluctuations and is reminiscent of the susceptibility of doped cuprates [121] and iron pnictide superconductors [122]. It will be interesting to test in future studies whether hole doping of $LaNiO_2$ alters the shape of $\chi_{\text{corr}}(T)$, similarly to cuprates where the broad maximum in the susceptibility shifts to lower temperatures with increasing doping concentration x and vanishes for heavy doping [121]. Moreover, the residual entropy detected in the specific heat C_p of $LaNiO_2$ (Fig. 1)

is compatible with the presence of paramagnons, which are a hallmark feature of doped cuprates. This suggests a possible analogy between the parent (undoped) IL nickelates and doped cuprates.

A good starting point to understand “nonlocal spin fluctuations” from a wave-function perspective can be models in the localized limit, such as the Heisenberg model or finite-size clusters. As shown in Fig. 6 already a simple 2×2 cluster (and, even more accurately, an embedded cluster via CDMFT) can give rise to broad maxima and deviations from Curie-Weiss behavior in their static susceptibility without symmetry breaking. This is due to the formation of intersite singlet ground states and thermal occupation of spin triplets. Another well-known singlet mechanism that leads to a maximum in the static susceptibility is found in the Kondo model where a fully localized magnetic impurity hybridizes with delocalized conduction electrons [136]. The ansatz to interpret the $3d$ electrons of Ni as fully localized and oxygen $2p$ or rare-earth $5d$ states as a delocalized bath was addressed in Refs. [28,34]. However, when sizable $3d$ - $3d$ hoppings are included in realistic *ab initio* material models, rather a Hubbard-type scenario is supported. In either case the mentioned singlets are nonlocal non-single-Slater-determinant states. In order to capture their impact on the susceptibility we can therefore neither resort to effective single-particle (like DFT) nor to purely local theories (like DMFT). On the other hand, DGA not only includes short-range spin fluctuations, such as the mentioned singlets, but fluctuations on all length scales.

Importantly, we find that our DGA calculations (Fig. 6) qualitatively reproduce the salient temperature dependence of the experimental χ_{corr} in Fig. 5. Therewith, an interpretation of the peculiar susceptibility $LaNiO_2$ is provided since in the framework of DGA the occurrence of a maximum in χ^{DGA} and a substantial downturn at low temperatures can be attributed to an increasing influence of nonlocal fluctuations. Furthermore, this lends support to the notion that strong magnetic fluctuations suppress the ordering tendencies in parent IL nickelates and prevent the formation of long-range magnetic order down to lowest temperatures. At a more fundamental level, long-range order is not occurring in our calculations and not accompanying any strong fluctuations at finite temperatures due to the fact that DGA in two dimensions obeys the Mermin-Wagner theorem. Note that this is in contrast to DMFT calculations, where finite-temperature ordering is possible due to its mean-field nature with an ordering temperature of $T_{\text{Néel}}^{\text{DMFT}} \approx 1500$ K. In cuprates, long-range AFM order emerges in spite of a quasi-2D character of the lattice and electronic structure due to anisotropic exchange generating an Ising-type component of the order parameter in compounds with frustrated interlayer exchange [137], or unfrustrated 3D coupling [138]. The absence of apical oxygen and vanishing hybridization between Ni $3d$ and O $2p$ states [28] might hamper analogous couplings in IL nickelates. The introduction of a hypothetical hopping in the c direction in IL nickelates would lift the 2D constraint of the Mermin-Wagner theorem also for DGA and we would expect to obtain a long-range ordered ground state. However, the long-range order would set in at lower temperatures with respect to DMFT due to the additional consideration of nonlocal fluctuations [92,93].

A comparison of Figs. 6 and 5 shows that the broad maximum in $\chi^{\text{D}\Gamma\text{A}}$ around $T_*^{\text{D}\Gamma\text{A}} \approx 900$ K is compatible with the increase of the experimental χ_{corr} with an onset above 65 K. Note that the low-temperature upturn below 65 K is likely related to paramagnetic impurities, which are not included in our theoretical modeling. The compatibility above 65 K strengthens the conjecture that the materials class of IL nickelates is a close realization of the single-band Hubbard model [19]. Physically, this remarkable degree of applicability might be rooted in the absence of Zhang-Rice singlets [28], due to the exceptionally large energetic distance of the oxygen degrees of freedom in IL nickelates [27,28]. However, as already stated in the Results section, the magnitude of the theoretically calculated susceptibility is significantly smaller than the experimental determined values. Notably, the different employed computation methods all yield susceptibilities of the same order of magnitude. Thus, the discrepancy to experiment might be explained by our initial assumption for the model, i.e., a half-filled Hubbard model with a pure Ni $3d^9$ configuration. Additional D Γ A studies exploring multi-band effects [139] and a variation of U are highly desired for addressing this issue. Furthermore, it cannot be ruled out that χ_{corr} contains contributions beyond such obvious considerations. For instance, it was proposed that topotactically reduced IL nickelates are prone to the inclusion of hydrogen [140]. The resulting high-spin Ni $3d^8$ contributions, not captured by our model, could enhance the magnetic response substantially. Similarly, the aforementioned remaining apical oxygen due to incomplete reduction can induce local $3d^8$ -like configurations. It is possible that these isolated Ni $^{2+}$ spins are exchange coupled to the square-planar Ni $^{1+}$ network and therefore exhibit the same temperature-dependent susceptibility, albeit with a larger amplitude.

Along the lines of a possible analogy between undoped IL nickelates and doped cuprates, an evident question is whether the former material class exhibits a pseudogap similar to that of lightly doped cuprates. More specifically, in cuprates, the hallmark for the original identification of the pseudogap was the suppression of the nuclear magnetic resonance (NMR) Knight shift [141]. In linear response calculations this suppression corresponds to the formation of a maximum in the uniform magnetic susceptibility [60,88]. Hence, our complementary CDMFT and D Γ A analysis indicates that a pseudogaplike regime emerges below $T_*^{\text{D}\Gamma\text{A}} \approx 900$ K and the observed non-Curie-Weiss behavior of $\chi_{\text{corr}}(T)$ is compatible with this notion. Notably, a pseudogap temperature T_* higher than 400 K is also consistent with a recent ^{139}La NMR study on polycrystalline LaNiO_2 [22]. Assuming that our modelization of LaNiO_2 as a single-band Hubbard model is appropriate, the opening of this pseudogap can be attributed to nonlocal magnetic correlations [60,142,143].

Remarkably, recent RIXS experiments determined the AFM exchange coupling J to be as large as ~ 65 meV in NdNiO_2 [23], which is in good agreement with 77 meV obtained from many-body quantum chemistry calculations [12] and reminiscent of the strong exchange of up to ~ 180 meV in IL cuprates [144]. In a simplified picture, considering the strong coupling limit, our choice of parameters corresponds to an exchange coupling of $J = 4t^2/U \sim 198$ meV. However, such mapping of our model in the localized limit of the

Heisenberg model and strong coupling limit is likely not appropriate and rather suggests that the coupling is substantially reduced.

V. CONCLUSION

In summary, we presented a comprehensive study of the magnetic correlations in LaNiO_2 combining experiment and quantum many-body theories. The observed signatures of strong nonlocal spin fluctuations, paramagnons, and a pseudogap suggest a striking analogy between the parent IL nickelates and doped cuprates.

Taking these parallels one step further, the question arises as to whether the long-range AFM phase of parent cuprates can be invoked also in IL nickelates, for instance, by electron doping or tuning of the effective bandwidth as well as the Ni-O hybridization. More generally, future multimethod studies using the complementary techniques DMFT, CDMFT, and D Γ A can explore how charge-carrier doping affects the magnetic correlations present in parent IL nickelates [145] and in particular to what degree they persist at doping levels that host a superconducting ground state. Moreover, an investigation of the rare-earth series RNiO_2 can provide insights on whether hybridization with the rare-earth $5d$ bands leads to a quantitative deviation from the qualitative applicability of the single-band Hubbard model, which we demonstrated for LaNiO_2 .

Finally, our study lends support to the notion that AFM spin fluctuations are a prime candidate for mediating superconductivity in IL nickelates.

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APPENDIX A: CRYSTAL STRUCTURE AND STOICHIOMETRY

Powder x-ray diffraction (PXRd) of LaNiO_3 and the reduced sample (LaNiO_2) was performed at room temperature using a Rigaku Miniflex diffractometer with $\text{Cu } K_\alpha$ radiation (Fig. 7). Rietveld refinements were conducted with the FULL-

TABLE I. (a) Refined atomic coordinates of LaNiO_3 in rhombohedral space group $R\bar{3}c$ (hexagonal axes) extracted from powder x-ray diffraction measurements (Fig. 7). (b) Refined atomic coordinates of LaNiO_2 in tetragonal space group $P4/mmm$.

(a) LaNiO_3 $a, b = 5.45095(4) \text{ \AA}, c = 13.13313(11) \text{ \AA}$				
Atom	x	y	z	$U(\text{\AA}^2)$
La (6a)	0	0	0.25	1.097(8)
Ni (6b)	0	0	0	0.674(17)
O (18e)	0.54430(62)	0	0.25	1.637(64)
Reliability factors	χ^2	R_B	R_f	R_{wp}
	7.53(2)	6.67	6.22	12.7
(b) LaNiO_2 $a, b = 3.95496(4) \text{ \AA}, c = 3.36582(5) \text{ \AA}$				
Atom	x	y	z	$U(\text{\AA}^2)$
La (1d)	0.5	0.5	0.5	0.644(10)
Ni (1a)	0	0	0	0.494(31)
O (2f)	0	0.5	0	0.039(67)
Reliability factors	χ^2	R_B	R_f	R_{wp}
	6.61(2)	7.34	5.06	21.0

PROF software suite [146]. LaNiO_3 and LaNiO_2 were refined in rhombohedral space group $R\bar{3}c$ and tetragonal $P4/mmm$,

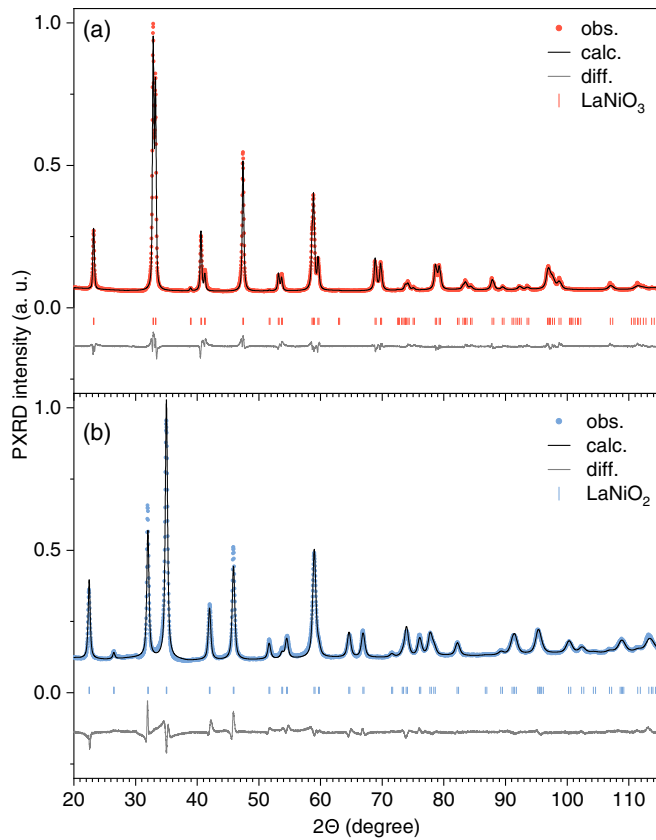


FIG. 7. Powder x-ray diffraction pattern of LaNiO_3 (a) and LaNiO_2 (b) at $T = 300 \text{ K}$ measured with $\text{CuK}\alpha$ radiation. Solid black lines correspond to the calculated intensities from the Rietveld refinements. The calculated Bragg peak positions are indicated by vertical bars and the differences between the experimental and calculated intensities are shown as solid grey lines at the bottom.

respectively. The refined structural parameters are presented in Table I. PXRD did not indicate the presence of any impurity phases. We note, however, that our laboratory-based standard PXRD measurement does not allow for ruling out subtle deviations from the ideal $P4/mmm$ structure, such as a buckling of the Ni-O bonds in the NiO_2 planes, which would imply a deviation from the 180° O-Ni-O bond angles. Similarly to Refs. [47–49], we observe anisotropic line-shape broadenings in our PXRD data, leading to deviations in the Rietveld analysis and increased R factors.

The stoichiometry of the samples was determined using scanning electron microscopy (SEM) with energy-dispersive x-ray spectroscopy (EDS) (Fig. 8) and inductively

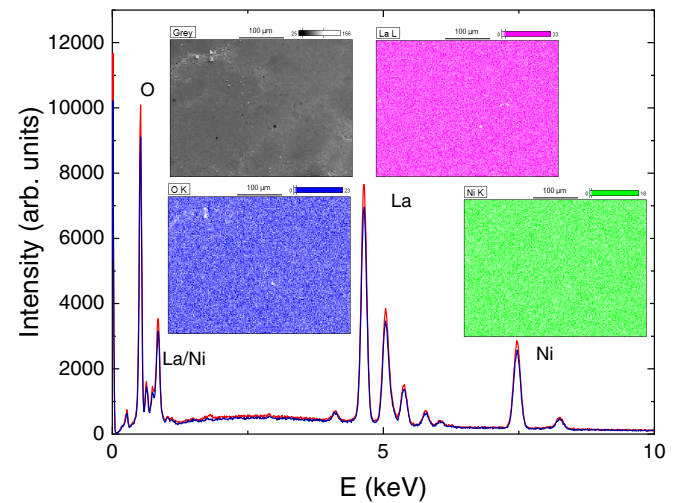


FIG. 8. Energy dispersive x-ray spectra (EDS) of a pellet of LaNiO_2 . The red line corresponds to an area scan, while the blue line is from a local point. The insets show the EDS maps of different elements, with Ni in green, La in magenta, and O in blue. No sudden jumps in Ni compared to La were observed.

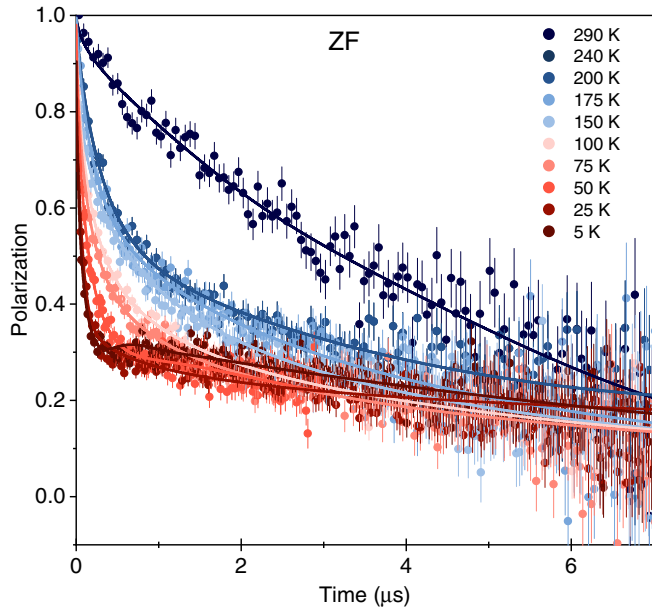


FIG. 9. ZF μ SR spectra measured at various temperatures. Solid lines are fits to the data (see text).

coupled plasma mass spectroscopy (ICP-MS). The latter method indicates the stoichiometries $\text{La}_{0.99(1)}\text{Ni}_{0.99(1)}\text{O}_{2.99(3)}$ and $\text{La}_{0.99(1)}\text{Ni}_{0.99(1)}\text{O}_{2.02(3)}$, corresponding to the ideal stoichiometries (within the experimental error) of the perovskite and infinite-layer phase, respectively. The former method indicates that a pressed LaNiO_2 pellet exhibits a highly homogeneous distribution of all elements down to the resolution limit (see inset in Fig. 8). Together with PXRD, we can estimate that ferromagnetic Ni inclusions in the LaNiO_2 sample are below 2.5 wt.%.

APPENDIX B: COMPLEMENTARY μ SR DATA

Figure 9 shows the full set of ZF μ SR spectra measured at various temperatures. Figure 10 displays μ SR spectra measured at 290 K in different longitudinal fields (LF). The indicated fits with a dynamical Lorentzian Kubo-Toyabe model describe the data reasonably well, whereas a static model is inconsistent with the data (not shown here). From the fit we obtain a fluctuation rate of approximately 0.5 MHz at 290 K.

APPENDIX C: COMPLEMENTARY SUSCEPTIBILITY DATA

Figure 11(a) shows the temperature dependence of the susceptibility χ of LaNiO_2 in different external magnetic fields. As expected for a spin glass, upon increasing field, the cusp in the ZFC curves shifts to lower temperatures and the ZFC and FC curves join together at lower temperatures.

We identify two different regimes where the irreversibility of the ZFC and FC curves is present [Fig. 3(a)]. In the low-temperature regime around T_f and below, the width of the hysteresis in the isothermal magnetic hysteresis loops is particularly broad [Fig. 11(b)] and we attribute the ZFC-FC splitting mostly to the glassy character of the system. In gen-

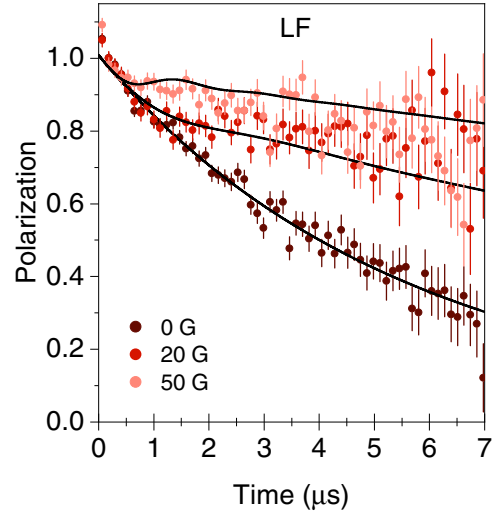


FIG. 10. μ SR spectra at 290 K in different longitudinal fields (LF). Solid black lines are dynamical fits (see text).

eral, spin glasses can exhibit a remnant magnetization in the temperature region of the freezing transition and below, due to the irreversible behavior of the magnetization. This remnant magnetization depends in a detailed way on the “magnetic history” of the sample [114] and evolves with time; i.e., it does not reflect the thermal equilibrium behavior of the glass. Such hysteretic effects can occur not only in spin glasses with underlying ferromagnetic correlations, but also in the antiferromagnetic case, thus precluding such distinction in our measurement. In antiferromagnetic glasses, hysteresis can be observed in particular in case of slow dynamics and/or uncompensated spins at surfaces of clusters.

The persisting ZFC-FC splitting in the regime above T_f in Fig. 11(a) is possibly induced by small amounts of ferromagnetic impurities, such as elemental Ni. This splitting is relatively narrow and also the widths of the hysteresis loops decrease significantly above T_f [Fig. 11(b)]. Nevertheless, it is also possible that the observed ZFC-FC splittings and hysteresis do not correspond to two different regimes, but

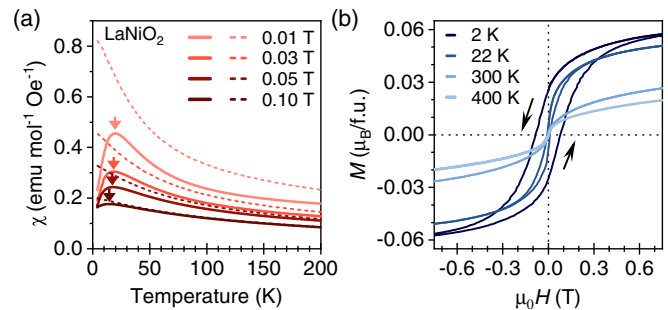


FIG. 11. (a) Static (dc) magnetic susceptibility χ of LaNiO_2 in different external magnetic fields measured upon heating after zero-field cooling (ZFC, solid lines) and field cooling (FC, dashed lines), respectively. The arrows indicate the positions of the cusps in the ZFC curves. (b) Isothermal magnetic hysteresis loops of LaNiO_2 measured up to $\mu_0 H = 0.75$ T at different temperatures. Black arrows indicate the directions of the field sweep.

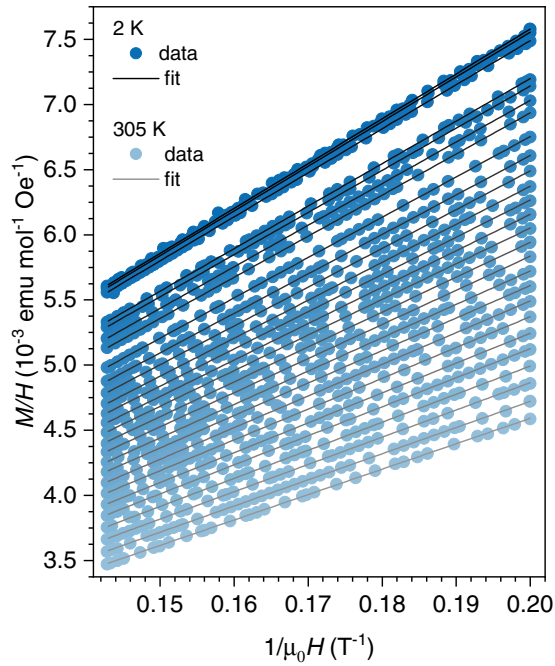


FIG. 12. Isothermal magnetization of LaNiO₂ measured between 5 and 7 T at 2, 4, 6, 8, 10, 15, 20, 35, 50, 65, 80, 95, 110, 125, 140, 155, 170, 185, 200, 215, 230, 245, 260, 275, 290, and 305 K, respectively (filled symbols from dark blue to light blue). The data (M/H) are plotted as a function of $1/H$, according to the Honda-Owen method. Solid lines (from dark to light gray) are linear fits for each temperature. The y -intercept of each linear fit corresponds to the corrected paramagnetic susceptibility χ_{corr} displayed in Fig. 5 in the main text.

are a consequence of either the spin glass or Ni impurities. Moreover, a coupling between the two phenomena could be responsible for the enhanced width of the hysteresis at low temperatures.

Note that our sample characterization (see Methods section and Appendix A) indicates that the amount of (ferromagnetic) impurities is very small. This notion is supported by the magnetization measurements shown in Fig. 11(b), revealing that the saturation magnetization in units of μ_B/Ni is low. In particular, the highest observed magnetization values are still substantially smaller than the expected value of $0.6\mu_B$ for metallic Ni, ruling out the presence of a bulk ferromagnetic Ni phase. Hence, we conclude that the Ni impurity contribution in our sample is significantly less than 2.5 wt.%.

APPENDIX D: CORRECTED SUSCEPTIBILITY AND MODEL FITS

Figure 12 shows the raw data of the isotherms of LaNiO₂ together with the fits from which the corrected susceptibility $\chi_{\text{corr}}(T)$ in Fig. 5 of the main text was extracted. We used the Honda-Owen method to determine the intrinsic susceptibility from the magnetization isotherms measured at high magnetic fields. We consider two contributions in the measured magnetization: one originating from the intrinsic paramagnetic susceptibility and the other from ferromagnetic impurities,

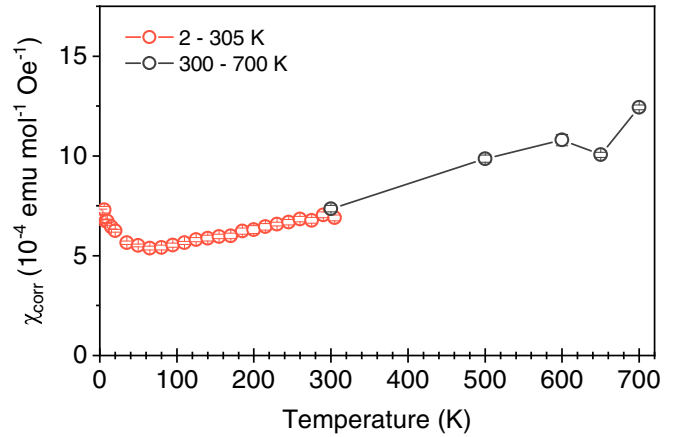


FIG. 13. Corrected paramagnetic susceptibility χ_{corr} of LaNiO₂ across a wide temperature range. Red data points are reproduced from Fig. 5 in the main text. Dark gray data points were extracted from isotherms (Fig. 14) measured in a high-temperature setup (see Methods).

corresponding to

$$M = \chi_{\text{corr}}H + C_{\text{sat}}M_{\text{sat}}, \quad (\text{D1})$$

where χ_{corr} is the intrinsic susceptibility, H is the applied magnetic field, and M_{sat} is the magnetization due to impurities. Then, in the limit $\mu_0H \rightarrow \infty$, the y -intercept in a M/H vs $1/\mu_0H$ plot corresponds to the intrinsic susceptibility χ_{corr} (see Fig. 12). We note that in Ref. [48] a method (“subtraction method”) closely similar to the Honda-Owen method was applied to extract the susceptibility of polycrystalline LaNiO₂, however, from magnetization isotherms measured only up to 5 T.

In addition to the data shown in Fig. 12, isotherms between 5 and 7 T were measured at elevated temperatures and the magnetic susceptibility was extracted. As can be seen in Figs. 13 and 14, the monotonous increase of $\chi_{\text{corr}}(T)$

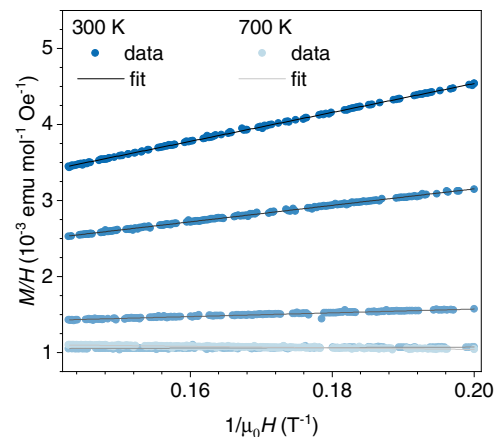


FIG. 14. Isothermal magnetization measured between 5 and 7 T at 300, 500, 600, 650, and 700 K, respectively (filled symbols from dark blue to light blue). Solid lines (from dark to light gray) are linear fits for each temperature to extract χ_{corr} (see Fig. 13). The data (M/H) are plotted as a function of $1/\mu_0H$, according to the Honda-Owen method.

of LaNiO₂ continues at least up to 600 K. At even higher temperatures, the small dip in the curve around 650 K is likely associated with the presence of Ni impurities, which are estimated to be significantly less than 2.5 wt.% in our pristine LaNiO₂ sample (see Appendix C) while additional impurities might be created during the heating process. These Ni impurities apparently have a small effect on $\chi_{\text{corr}}(T)$, even though χ_{corr} is extracted via the Honda-Owen method from high-field isotherms between 5 and 7 T. Note that the increase after 650 K in signal is due to the oxidization and decomposition of the sample as revealed by repeated measurement and PXRD analysis.

We conclude that the temperature dependence of χ_{corr} of our LaNiO₂ sample well below 600 K is not qualitatively dominated by the Ni impurities. Specifically, we note that the temperature dependence of the magnetic Knight shift in a recent ¹³⁹La NMR study on polycrystalline LaNiO₂ [22] is qualitatively consistent with our $\chi_{\text{corr}}(T)$ in Fig. 13, indicating that our $\chi_{\text{corr}}(T)$ essentially reflects the intrinsic susceptibility of LaNiO₂ and not that of Ni impurities.

APPENDIX E: COMPUTATIONAL DETAILS OF THE NUMERICAL CALCULATIONS

1. DMFT and DGA

We apply DGA in its ladder version with Moriyasque λ corrections in the spin channel [62,63,83] since it is particularly suited for the description of low-dimensional systems with strong nonlocal magnetic fluctuations in the paramagnetic phase (paramagnons) [62,80,147,148] due to its incorporation of nonlocal ladders in the relevant particle-hole channels [62]. We solve the Bethe-Salpeter equations in Matsubara frequency space with a maximum of $N_{i\omega}=100$ positive fermionic and $N_{i\Omega}=99$ positive bosonic Matsubara

frequencies for the two-particle Green function at the lowest temperature shown. We extrapolated the physical susceptibility in the number of fermionic Matsubara frequencies $N_{i\omega} \rightarrow \infty$ according to $\chi = a + b/N_{i\omega}^2$. We used 128 linear momentum grid points. In our calculations the dc susceptibility corresponds to the zeroth bosonic Matsubara frequency and zero momentum transfer, i.e., $\chi^{\text{DGA}} \equiv \text{Re } \chi_m(\mathbf{q}=(0,0), i\Omega_n=0)$. For the solution of the self-consistently determined Anderson impurity model we used the continuous-time quantum Monte Carlo solver in its interaction expansion CT-INT [149–151] as part of the TRIQS [99] package. We used 256×10^5 cycles and 6200 core hours per temperature point.

2. CDMFT

In order to calculate the magnetic susceptibility in CDMFT, analogously to experiment, we applied a small ferromagnetic (uniform) field with strength $H_F = 0.008$ and measured the mean uniform magnetization m of the cluster:

$$\text{Re } \chi_m(\mathbf{q}=(0,0), i\Omega_n=0) = \left. \frac{\partial m}{\partial H} \right|_{H=0} \approx \frac{m}{H_F}. \quad (\text{E1})$$

We checked that this value of the applied field is within the linear response regime. We again used the continuous-time quantum Monte Carlo solver in its interaction expansion CT-INT as part of the TRIQS package. We converged the calculations using 30 iterations with a statistic of 256×10^6 cycles for each iteration and 4000 total core hours per temperature point.

3. Unit conversion

We converted the computed susceptibilities given in units of ($\mu_B^2 \text{ eV}^{-1}$) to ($\text{emu mol}^{-1} \text{ Oe}^{-1}$) by multiplying with the factor 3.233×10^{-5} ($\text{emu mol}^{-1} \text{ Oe}^{-1} \mu_B^{-2} \text{ eV}$).

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