Precision search for magnetic order in the pseudogap regime of $La_{2-x}Sr_xCuO_4$ by muon spin relaxation

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We report a high-precision search for orbital-like magnetic order in the pseudogap region of $La_{2-x}Sr_xCuO_4$ single crystals using zero-field muon spin relaxation (ZF- μ SR). In contrast to previous studies of this kind, the effects of the dipolar and quadrupolar interactions of the muon with nearby nuclei are calculated. ZF- μ SR spectra with a high number of counts were also recorded to determine whether a magnetically ordered phase exists in dilute regions of the sample. Despite these efforts, we find no evidence for static magnetic order of any kind in the pseudogap region above the hole-doping concentration p = 0.13.

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I. INTRODUCTION

A distinctive feature of high-transition-temperature (T_c) cuprate superconductors is the pseudogap region that exists above T_c and over a wide range of doping. For some time there has been much debate on whether the pseudogap is a manifestation of a phase transition. Recently, it has been demonstrated that the onset of the pseudogap in optimally doped Pb_{0.55}Bi_{1.5}Sr_{1.6}La_{0.4}CuO_{6+ δ} at a temperature $T^* \approx$ $3.5T_c$ is likely a sign of a phase transition to a nonsuperconducting state with a broken symmetry.¹ Although this is not necessarily a magnetically ordered state, proposed orders for the pseudogap state include time-reversal-symmetry-breaking phases that contain ordered circulating orbital currents, which either break^{2,3} or preserve⁴ translational symmetry. The strongest experimental evidence for an orbital-current phase are the observations of an unusual translational-symmetry preserving magnetic order in YBa₂Cu₃O_y and HgBa₂CuO_{4+ δ} by spin-polarized neutron diffraction,⁵⁻⁹ which bear some resemblance to the ordered Θ_{II} circulating-current phase proposed in Ref. 4. It is worth pointing out, however, that by extrapolation the onset of this orbital-like magnetic order is expected to occur near T_c at optimal doping, and hence its relationship to the phase transition reported in Ref. 1 is unclear. Furthermore, to date such orbital-like magnetic order has not been observed in any cuprate beyond a hole doping of p = 0.135, whereas the Θ_{II} phase is predicted to persist up to $p \sim 0.19$.

In contrast to the experimental techniques used in the above studies, local probes of magnetism, such as nuclear magnetic resonance (NMR) and zero-field muon spin relaxation (ZF μ SR) provide information on the magnetic volume fraction. Unfortunately, ⁸⁹Y NMR experiments on Y₂Ba₄Cu₇O_{15- δ},¹⁰ Zeeman perturbed nuclear quadrupole resonance (NQR) measurements of YBa₂Cu₄O₈,¹¹ and ZF- μ SR experiments on YBa₂Cu₃O_y (Refs. 12 and 13) and La_{2-x}Sr_xCuO₄ (Ref. 14) have found no evidence for the onset of magnetic order at the pseudogap temperature *T**. One exception is the finding of anomalous magnetic order by ZF- μ SR¹³ in the same YBa₂Cu₃O_{6.6} single crystal studied in Ref. 6. The magnetic order is characterized by an onset temperature and an average local dipolar magnetic field that are in quantitative agreement

with the orbital-like magnetic order detected by polarized neutron diffraction. Yet the ZF- μ SR measurements clearly show this form of magnetic order existing in only about 3% of the sample, suggesting that it is associated with a minority phase in lower-quality samples.

Orbital-like magnetic order has also been observed in La_{1.915}Sr_{0.085}CuO₄ by polarized neutron diffraction,¹⁵ but is less pronounced than the long-range magnetic order that has been reported in the pseudogap regions of YBa₂Cu₃O_y (Refs. 5 and 6) and HgBa₂CuO_{4+ δ} (Ref. 7). In particular, the magnetic order in La_{1.915}Sr_{0.085}CuO₄ is short range, two (rather than three) dimensional, and occurs at a temperature far below *T*^{*}. Yet no such magnetic order was observed in a ZF- μ SR study of $x \ge 0.13$ samples by MacDougall *et al.*¹⁴ Nevertheless, the subtle nature of the magnetic order observed in La_{1.915}Sr_{0.085}CuO₄ warrants a more precise ZF- μ SR search to verify its existence and to determine whether it is an intrinsic property of the pseudogap phase of La_{2-x}Sr_xCuO₄.

Here we report two significant advances in the application of ZF- μ SR to search for magnetic order in the pseudogap region of La_{2-x}Sr_xCuO₄. First, we accurately determine the interactions of the positive muon (μ^+) with the nuclear spin system, allowing us to identify any residual relaxation of the ZF- μ SR spectrum that could be ascribed to static magnetic order. In doing so we have accurately identified the muon stopping site. Second, we have acquired ZF- μ SR spectra of higher statistics than in previous works, enabling a search for dilute or short-range magnetic order.

II. EXPERIMENTAL DETAILS

The experiments were performed on platelet-like single crystals of $La_{2-x}Sr_xCuO_4$ cut from a travelling-solvent floating zone (TSFZ) growth rod. The TSFZ growth procedure that was followed is described elsewhere.¹⁶ The single crystals cut from the TSFZ rod were annealed at 800 to 900 °C in an oxygen partial pressure to remove oxygen defects in accordance with the oxygen nonstoichiometry of $La_{2-x}Sr_xCuO_4$.¹⁷ Strontium concentrations greater than x = 0.125 were chosen to ensure the absence of static antiferromagnetism or the spin-glass–like magnetism previously observed by ZF- μ SR in lower-doped samples.^{18,19} Magnetic susceptibility measurements of the bulk superconducting transition temperature by a superconducting quantum interference device (SQUID) yield $T_c = 37.6, 37.3, 28$ and 17 K for the x = 0.15, 0.166, 0.216, and 0.24 samples, respectively.

The ZF- μ SR measurements of the La_{2-r}Sr_rCuO₄ single crystals were performed on the M15 and M20B surface muon beamlines at TRIUMF. Positive muons implanted into the sample Larmor precess about the local field B and decay according to $\mu^+ \rightarrow e^+ + \nu_e + \bar{\nu}_{\mu}$, with a mean lifetime $\tau \sim$ 2.2 μ s. The ZF- μ SR signal is generated from detection of the decay positrons, which are preferentially emitted along the muon spin direction. The samples were mounted with the crystallographic c axis of the $La_{2-x}Sr_xCuO_4$ single crystals parallel to the muon beam momentum. The initial muon spin polarization $\mathbf{P}(0)$ was oriented perpendicular to the c axis using a Wien filter. This has the advantage that neither positron detector directly faces the incoming muon beam. In this geometry the ZF- μ SR "asymmetry" spectrum is defined as the difference between the number of decay positrons sensed by scintillator detectors positioned above (A) and below (B)the sample, divided by the sum of the counts in these two detectors

$$\frac{N_A - N_B}{N_A + N_B} \equiv A(t) = a P(t), \tag{1}$$

where a < 1/3 is the initial asymmetry (dependent on the energy of the decay positrons and several experimental factors) and P(t) is the time evolution of the muon spin polarization. The latter is modeled by an appropriate relaxation function G(t):

$$P(t) = G(t)\cos(\gamma_{\mu}Bt), \qquad (2)$$

where γ_{μ} is the muon gyromagnetic ratio and *B* is the average internal magnetic field sensed by the muon. Note that B = 0 in the absence of magnetic order.

III. EXPERIMENTAL RESULTS

In previous studies of $La_{2-x}Sr_xCuO_4$ the contribution of the nuclear moments to the ZF- μ SR signal was assumed to be described by a static Gaussian Kubo-Toyabe (KT) relaxation function^{20,21}

$$G_{\rm KT}(t) = \frac{1}{3} + \frac{2}{3}(1 - \Delta^2 t^2) \exp\left[-\frac{1}{2}\Delta^2 t^2\right],$$
 (3)

where Δ is the second moment of the local magnetic field distribution at the muon site. The relaxation function $G_{\rm KT}(t)$ assumes that the muon is immobile, the local magnetic fields acting on the muon spin are static, and that the local magnetic field distribution is isotropic and Gaussian. The nuclear dipolar fields sensed by the positive muon μ^+ are usually static, because correlation times of the nuclear moments are generally much longer than the muon lifetime. Assuming that the nuclear dipolar fields acting on the muon spin are randomly oriented and that each muon sees a unique local field over the duration of its lifetime, 1/3 of the muon spins will be parallel to the field and not evolve in time. Consequently, the recovery of the muon spin polarization to 1/3 of its initial value at late times is characteristic of static fields. However, in a single crystal the contribution of the surrounding



FIG. 1. (Color online) ZF- μ SR signal of single crystal La_{1.784}Sr_{0.216}CuO₄ at T = 30 K recorded with the initial muon spin polarization **P**(0) perpendicular to the *c* axis. The solid green curve is a fit of the data below $t = 6 \ \mu$ s to Eq. (3) multiplied by an asymmetry factor *a*. The solid black curve is a fit to Eq. (4).

nuclear moments can substantially deviate from the random field approximation, with no recovery of P(t) to 1/3. For example, Fig. 1 shows that the static Gaussian Kubo-Toyabe function does not fully describe the ZF- μ SR spectrum of single crystal La_{1.784}Sr_{0.216}CuO₄ for the case where **P**(0) is perpendicular to the *c* axis—despite being in a region of the phase diagram where only the nuclear dipole moments are expected to contribute to the ZF relaxation function. Of particular note, the muon spin polarization dips below zero, whereas $G_{\text{KT}}(t)$ does not. To circumvent this problem the ZF- μ SR signals for La_{2-x}Sr_xCuO₄ in Ref. 14 were truncated at $t = 8 \ \mu$ s to facilitate fits to Eq. (3).

To more accurately portray changes in the functional form of the ZF- μ SR signal as a function of temperature and Sr content *x* (i.e., hole-doping concentration), here we fit the ZF- μ SR spectra to a simple phenomenological power-exponential relaxation function, such that

$$P(t) = \exp[-(\Lambda t)^{K}] \cos[\gamma_{\mu} B(T)t], \qquad (4)$$

with the relaxation rate Λ and the exponent *K* treated as temperature-independent quantities, and the average local magnetic field *B* being the only fit parameter allowed to vary with temperature. Figure 2(a) shows representative time spectra for La_{1.834}Sr_{0.166}CuO₄ that are well described by Eq. (4) over the entire 10 μ s time range. A fit to Eq. (4) is also shown for La_{1.784}Sr_{0.216}CuO₄ in Fig. 1. The evolution of the ZF- μ SR signal with temperature is reflected in the temperature dependence of *B* shown in Fig. 3(a). Consistent with the findings of MacDougall *et al.*,¹⁴ the ZF- μ SR signal does not exhibit a temperature dependence characteristic of a magnetic phase transition in any of the samples. At *T* = 200 K, the value of *B* is clearly reduced, but by the same amount at *x* = 0.15 and *x* = 0.24. Hence, the reduction of *B* at high temperatures is likely caused by muon diffusion, whereby the mobile μ^+ sees



FIG. 2. (Color online) (a) Comparison of the ZF- μ SR signals of La_{1.834}Sr_{0.166}CuO₄ at T = 2.5 K and T = 158 K measured over a 10 μ s time range with **P**(0) perpendicular to the *c* axis. (b) The early-time ZF- μ SR signal of La_{1.834}Sr_{0.166}CuO₄ at T = 2.5 K shown over the first 1 μ s. In both figures the solid curves are fits to the relaxation function of Eq. (4).

a time-averaged field over its lifetime. This same conclusion was reached in Ref. 14.

The dependence of Λ and K on the Sr content x is shown in Fig. 3(b). There is some increase in the values of both parameters with increasing x, but a clear reduction of Λ and K for the x = 0.24 sample. The former behavior may be the result of antiferromagnetic fluctuations, dilute regions of static magnetism and/or the relaxation caused by the Sr nuclei. While the precise source is unclear, there is no evidence for static magnetic order in any of these samples. The smaller values of Λ and K at x = 0.24 indicate another contribution to the ZF- μ SR signal. This is likely paramagnetic moments that are known to be present in heavily overdoped La_{2-x}Sr_xCuO₄ above $x \sim 0.20$.^{22–25} The onset of the Curie-like paramagnetism is the probable cause of the slightly larger values of B at x = 0.216 and x = 0.24 in Fig. 3(a).



FIG. 3. (Color online) Results of fits to Eq. (4). (a) Temperature dependence of *B* for all four samples. (b) Dependence of the relaxation rate Λ (solid squares) and the exponent *K* (open circles) on Sr content *x*. Note that both Λ and *K* are temperature-independent fit parameters.

In Ref. 13, three distinct magnetic components were observed in the ZF- μ SR signal of a large YBa₂Cu₃O_{6.6} single crystal. Neutron scattering experiments on this same single crystal show that the sample contains magnetic order associated with the impurity "green phase" Y₂BaCuO₅²⁶ and two additional kinds of unusual magnetic order.^{6,26} While one of the latter two magnetic components is manifested as a slow relaxing component that is also observed in higherquality samples, the other two forms of magnetic order are discernible as small-amplitude, rapidly damped oscillatory components in the early time range of the ZF- μ SR signal. The small amplitudes indicate that the magnetic orders are confined to small volume fractions of the YBa₂Cu₃O_{6.6} single crystal. However, Fig. 2(b) shows that there are no such oscillatory components in the early-time ZF- μ SR signal of La_{1.834}Sr_{0.166}CuO₄. We have also examined the early-time spectra of the other samples at various temperatures above T = 2.3 K and likewise find no evidence for any kind of short-range or dilute magnetic order.

Next we calculate the contribution of the nuclei to the polarization function P(t), in an effort to fully account for the observed ZF- μ SR spectrum.

IV. CALCULATION OF NUCLEAR-INDUCED RELAXATION

Here we describe a general numerical method for calculating the ZF relaxation function resulting from the dipolar magnetic and quadrupolar electrostatic interactions of the μ^+ with an arbitrary number of neighboring nuclei. We consider only the interactions of the nuclei with the muon, and ignore interactions amongst the nuclei themselves. This approximation is justified by the size of γ_{μ} , which is about an order of magnitude larger than the gyromagnetic ratios of the nuclei in La_{2-x}Sr_xCuO₄. We also assume that the electric field gradient (EFG) at each nuclear site is due to the Coulomb field of the unscreened positively charged muon and the nonsymmetric charge distribution of the crystal itself (i.e., the crystal EFG).

The interactions between the muon spin (S = 1/2) and N surrounding nuclei of spin I, as well as the effect of the crystal EFG, is described by the following Hamiltonian:^{27,28}

$$H = \sum_{j=1}^{N} \left(H_{j}^{D} + H_{\mu j}^{Q} + H_{o j}^{Q} \right),$$
(5)

where

$$H_j^D = \frac{\hbar^2 \gamma_\mu \gamma_j}{r_j^3} [\mathbf{S} \cdot \mathbf{I}_j - 3(\mathbf{S} \cdot \mathbf{n}_j)(\mathbf{I}_j \cdot \mathbf{n}_j)], \qquad (6)$$

$$H^{Q}_{\mu j} = \hbar \omega^{Q}_{\mu j} [(\mathbf{I}_{j} \cdot \mathbf{n}_{j}) \cdot (\mathbf{I}_{j} \cdot \mathbf{n}_{j}) - I(I+1)/3], \quad (7)$$

$$H_{oj}^{Q} = \frac{\hbar\omega_{oj}^{Q}}{2} \left[I_{jz}^{2} - \frac{1}{3} I_{j} (I_{j} + 1) + \frac{1}{6} \eta \left(I_{j+}^{2} + I_{j-}^{2} \right) \right].$$
(8)

In the above equations, the term H_i^D is the dipole-dipole interaction between the positive muon and the *j*th nucleus, H_{ui}^Q is associated with the quadrupolar energy of the nuclear spin I_j due to the EFG generated by the positive muon, \mathbf{n}_j is the unit vector pointing in the direction along the straight line that connects the muon to the *j*th nucleus located a distance r_i away, and γ_{μ} and γ_{j} are the gyromagnetic ratios of the muon and nuclei, respectively. The quadrupolar coupling constant $\omega_{\mu j}^Q$ is proportional to $1/r_j^3$.²⁹ The term H_{oj}^Q represents the quadrupolar energy of the nuclear spin due to the crystal EFG, with a quadrupolar coupling constant ω_{oi}^Q . The constant η is an asymmetry parameter which specifies the symmetry of the crystal EFG around the nucleus.²⁸ In our calculation, the values of the quadrupolar coupling ω_{oi}^{Q} and the asymmetry parameter η for La and Cu are taken from the literature. In particular, $\omega_{oj}^Q = 2\pi v_Q$ with $v_Q = 34.0$, 31.0, and 6.40 Hz for ⁶³Cu, ⁶⁵Cu, and ¹³⁹La, respectively.^{30,31} At Cu sites, $\eta = 0.03$,³² while at La sites, $\eta = 0.02$.³¹ In addition, the weighted averages of the two isotopes ⁶³Cu and ⁶⁵Cu are used for the gyromagnetic ratio γ_{Cu} and the nuclear quadrupole moment Q_{Cu} of the Cu nuclei.

Since the only stable isotope of Sr with nonzero spin is ⁸⁷Sr with a natural abundance of 7%, the Sr nuclei can be neglected in the calculations. We note that this is consistent with the study by MacDougall *et al.*,¹⁴ which showed that the relaxation rate of the ZF- μ SR signal of La_{2-x}Sr_xCuO₄ varies little in the range 0.13 $\leq x \leq 0.30$. Consequently, it



FIG. 4. (Color online) Muon D site (0.120, 0, 0.219) in the tetragonal unit cell of La₂CuO₄, which best describes the ZF- μ SR signal of La_{2-x}Sr_xCuO₄ at $x \ge 0.13$ (see Fig. 7). The coordinate values are in multiples of their respective lattice constants a = 3.80 Å, b = 3.80 Å, and c = 13.12 Å.³³ The small shaded rectangle visually emphasizes that the muon at the D site near the apical O resides on the *a*-*c* face of the unit cell. The lower left of the picture shows the 5 La nuclei and the 1 Cu nucleus included in the calculation of P(t) for this muon site (two of the La atoms are labeled 1 and 2 to identify them in the full crystal structure).

is sufficient to perform our calculations for a positive muon residing in a single crystal of the parent compound La_2CuO_4 .

With knowledge of the Hamiltonian [Eq. (5)], the time dependence of the muon spin polarization may be calculated from the density matrix of the spin system as

$$P(t) = \operatorname{Tr} \left\{ \rho(0) e^{iHt/\hbar} \left[\sigma_{\mu} \otimes \left(\bigotimes_{j=1}^{N} \mathbf{1}_{D_{j}} \right) \right] e^{-iHt/\hbar} \right\}.$$
(9)

Here $\sigma_{\mu} = \sigma_{\mu} \cdot \mathbf{P}(0) = \sigma_x \sin \theta \cos \beta + \sigma_y \sin \theta \sin \beta + \sigma_z \cos \theta$ is the projection of the muon spin along the direction of the initial polarization $\mathbf{P}(0)$, with θ and β being the polar and azimuthal angles, respectively, between $\mathbf{P}(0)$ and the crystallographic axes (see Fig. 4). Note that, from Eq. (8), the crystal EFG sets the quantization axis to be along the *c* axis of La_{2-x}Sr_xCuO₄. The initial density matrix is $\rho(0)$, $D_j = 2I_j + 1$ is the spin degeneracy for the *j*th nuclear spin, and $\mathbf{1}_n$ is the $n \times n$ identity matrix.

For a system in which the nuclear spins are randomly oriented, the initial density matrix is

$$\rho(0) = \frac{1}{D} (\mathbf{1}_2 + \sigma_\mu) \otimes \left(\bigotimes_{j=1}^N \mathbf{1}_{D_j} \right), \tag{10}$$

where

$$D = 2\prod_{j=1}^{N} D_j \tag{11}$$



FIG. 5. (Color online) (a) Approximation method calculations of the muon spin polarization function P(t) for different muon sites, with the initial muon spin polarization P(0) perpendicular to the *c* axis and oriented at 45° with respect to the *a* axis. The spatial coordinates of the muon sites and the nuclei included in each calculation are shown in Table I. The sites M₁, T₁, T₂, and U₁ were previously considered in Ref. 34. (b) P(t) for the same muon site, but with different nuclei used in the calculation (see Table I).

is the dimensionality of the Hamiltonian matrix. The time evolution of the muon spin polarization P(t) may be determined exactly by diagonalizing the $D \times D$ Hamiltonian. The main limitation of this approach is that it becomes very computationally expensive as the number of nuclei (and hence D) increases. In this work the largest systems studied include 10 nuclei, corresponding to $D = 2^{26} \approx 6.71 \times 10^7$. We found that for these large values of D, it was necessary to use an approximate method to determine P(t), and hence we used a method developed by Celio to study P(t) in copper.²⁷ This approximation method is based on the Trotter formula and utilizes the random phase approximation. We verified good agreement with the exact calculation for some of our calculations with less nuclei.

The calculation of P(t) is strongly dependent on the orientation of P(0) with respect to the *c* axis, the number and type of nuclei, and the precise muon site. With the orientation of P(0) fixed in the experiment, we used the approximation method to perform an exhaustive search for the function P(t) that best describes the ZF- μ SR signal of La_{2-x}Sr_xCuO₄ using numerous combinations of the muon site and the surrounding

TABLE I. Muon sites of polarization functions plotted in Fig. 5. The spatial coordinates of the muon site are denoted by multiples of the lattice constants a, b, and c of La₂CuO₄. The nearest-neighbor nuclei used in the calculation of P(t) for each muon site are listed in the third column. Two Cu nuclei are denoted as 2Cu, a La nucleus at site 1 (see Fig. 4) is denoted by La(1), etc.

Label	Muon site	Nuclei
M	(0.5, 0.0, 0.096)	2Cu, 2La(2)
T1	(0.2, 0.0, 0.15)	2Cu, La(1), 2La(2)
T2	(0.225, 0.0, 0.225)	2Cu, La(1), 2La(2)
U1	(0.12, 0.0, 0.11)	2Cu, La(1), 2La(2)
А	(0.0, 0.0, 0.212)	5Cu, La(1), 4La(2)
Ba	(0.1, 0.0, 0.1)	Cu, La(1), 2La(2)
B _b	(0.1, 0.0, 0.1)	4Cu, La(1), 2La(2)
D	(0.12, 0.0, 0.219)	Cu, La(1), 4La(2)

nuclei. Figure 5(a) shows P(t) calculated by the approximation method for a handful of the numerous potential muon sites considered in our study. Some of these sites were considered in earlier μ SR studies.^{34,35} As demonstrated in Fig. 5(b), only nearest-neighbor nuclei need to be included in the calculation of P(t). In particular, note that both B_a and B_b correspond to the same muon site, but the calculation of P(t) for B_b includes three additional Cu nuclei located further away from the muon (see Table I). The strong dependence of P(t) on the muon site means that ZF- μ SR can be used to accurately determine the location of the implanted μ^+ in situations where there is a single muon site, the muon does not diffuse, and there are no additional sources of relaxation (e.g., electronic moments and/or loop-current order).

Of all the muon sites we considered, only the polarization function P(t) calculated for the site (0.120, 0, 0.219) shown in Fig. 4 (which we refer to here as the D site) accounts for the observed ZF- μ SR spectra (see Fig. 7). This site is located



FIG. 6. (Color online) Time evolution of muon spin polarization P(t) calculated for muon D site of Fig. 4 and different orientations of the initial muon spin polarization P(0). The upper dashed curve shows P(t) for P(0) parallel to the *c* axis. The lower solid curves show P(t) for different orientations of P(0) in the *a*-*b* plane, labeled by the angle between P(0) and the *a* axis.



FIG. 7. (Color online) Comparisons between P(t) calculated for muon A and D sites and (a) ZF- μ SR signal of La_{1.834}Sr_{0.166}CuO₄ at T = 158 K measured with **P**(0) perpendicular to the *c* axis, and (b) ZF- μ SR signals of La_{1.87}Sr_{0.13}CuO₄ at T = 144 K from Ref. 14 for **P**(0) perpendicular to the *c* axis and **P**(0) parallel to the *c* axis. Note that the calculation for **P**(0) \perp **c** assumes that the direction of **P**(0) makes an angle of 45° with the *a* axis. Also, the ZF- μ SR signals shown here have been divided by the initial asymmetry *a*.

approximately 0.7 Å from the apical oxygen, which agrees with the widely held view that the μ^+ bonds to an oxygen atom in cuprates. While it is not exactly one of the muon sites suggested in earlier works,^{34,35} it is consistent with ZF- μ SR measurements of the antiferromagnetic phase of La₂CuO₄ by Hitti *et al.*³⁵ that restrict the muon site to the *a-c* plane.

As shown in Fig. 4 and indicated in Table I, five La nuclei of spin I = 7/2 and one Cu nucleus of spin I = 3/2 are included in the calculation of P(t) for the muon D site. Figure 6 shows how P(t) is dependent on the orientation of P(0) with respect to the crystal lattice. Although the orientation of P(0) in the *a-b* plane was random in our experiments, the ZF- μ SR spectra were reproducible when the sample was rotated about the *c* axis to a different position. We can understand this as being a

consequence of the equivalency of the sites (0.120, 0, 0.219)and (0, 0.120, 219) in the tetragonal unit cell. Since these are occupied by the muon with equal probability, the ZF- μ SR spectrum is an average of P(t) for these two sites. For the case $\mathbf{P}(0) \perp \mathbf{c}$, at any orientation this superposition is equivalent to P(t) calculated with **P**(0) lying in the *a*-*b* plane and making an angle of 45° with the *a* axis. As shown in Fig. 7(a), there is fairly good agreement between the calculation assuming the muon is located at the D site and the ZF- μ SR spectrum of La_{1.834}Sr_{0.166}CuO₄ measured with $P(0) \perp c$. This is also shown to be the case for the A site, where the muon is positioned directly above the apical oxygen. To distinguish between these two sites we consider the dependence on the angle between P(0) and the *c* axis. In Fig. 7(b) the calculations of P(t) for both $\mathbf{P}(0) \perp \mathbf{c}$ and $\mathbf{P}(0) \parallel \mathbf{c}$ are compared to ZF- μ SR spectra of La_{1.87}Sr_{0.13}CuO₄ from Ref. 14. While these data are less accurate at late times due to lower muon counts, it is clear that the calculation for the A site fails to describe the ZF- μ SR signal with **P**(0) || **c**.

V. DISCUSSION AND CONCLUSIONS

The ZF- μ SR measurements presented here and in Ref. 14 show no evidence for any form of static magnetism that can be directly linked to the pseudogap phase of $La_{2-r}Sr_rCuO_4$. Our determination of the muon stopping site enables a good estimate of the magnitude of the average local magnetic field that should have been detected if static orbital-like magnetic order of the kind observed by spin-polarized neutron diffraction^{5-9,15} were present. The neutron results are in general agreement with the loop-current pattern of the Θ_{II} state proposed in Refs. 4 and 36, which consists of two oppositely circulating current loops per unit cell. If the orbital currents flow in the CuO₂ layers along the Cu-O and O-O bonds as originally proposed, a μ^+ residing at the D site (0.120, 0, 0.219) will experience a local field of 308 G/ μ_B . For the maximum possible ordered magnetic moment of 0.02 μ_B deduced from the spin-polarized neutron measurements of $La_{1.915}Sr_{0.085}CuO_4$,¹⁵ the corresponding dipolar magnetic field sensed by the muon is 6.2 G. In this case the damped ZF- μ SR signal should oscillate with a period of $\sim 12 \ \mu s$, such that one nearly complete oscillation is observed over the time range of our measurements.

The moments of the orbital-like magnetic order observed in hole-doped cuprates by spin-polarized neutron diffraction are actually pointing at an angle of roughly 45° with respect to the c axis. This is compatible with the orbital currents of the Θ_{II} phase flowing out and back into the CuO₂ layers through the apical oxygen atoms.³⁷ In this scenario, the corresponding moments that are perpendicular to the faces of the CuO₆ octahedra in $La_{2-x}Sr_xCuO_4$ point along directions making an angle of 61° away from the c axis. For this arrangement with an ordered moment of $0.02 \mu_B$, the average local field that a muon at the D site in $La_{2-x}Sr_xCuO_4$ would detect is about 90 G. Since 0.02 μ_B is the maximum possible value of the ordered moment observed in La_{1,915}Sr_{0.085}CuO₄, and the orbital-like ordered moment in other cuprates^{5–9} generally decreases with increased hole-doping, one might think of 90 G as an upper limit for the average local field in the $x \ge 0.13$ samples considered here. On the other hand, a larger value is possible if the ordered moment in La_{1.915}Sr_{0.085}CuO₄ is reduced due to a higher degree of disorder and/or competition of the orbital-like magnetic order with the spin-density wave (SDW) order present in lower-doped samples. Competition with SDW order has been suggested as a potential explanation for the smaller ordered moment observed in YBa₂Cu₃O_{6.45} compared to that of YBa₂Cu₃O_{6.5}.⁸ Regardless, the contribution of orbital-like magnetic order to the ZF- μ SR signal will be strongly damped if it is short range as in the case of La_{1.915}Sr_{0.085}CuO₄. Yet no such component is observed in any of the La_{2-x}Sr_xCuO₄ samples we measured.

The failure here and in Ref. 14 to detect orbital-like magnetic order in $La_{2-x}Sr_xCuO_4$ of the kind observed by spin-polarized neutron diffraction may indicate that the local fields are rapidly fluctuating outside the μ SR time window. It has also been suggested that the μ^+ may destroy loopcurrent order in cuprates,³⁸ although thus far it has also eluded detection by NMR or NQR. On the other hand, since the polarized neutron diffraction experiments cannot deduce magnetic volume fractions, the orbital-like magnetic order could still be associated with a small minority phase that evolves with hole doping.¹³ Having said all of this, there is currently no disagreement between the polarized neutron diffraction and ZF- μ SR experiments on $La_{2-x}Sr_xCuO_4$, since there is no overlap in the doping range of the samples studied by these two techniques. The ZF- μ SR experiments have focused on $x \ge 0.13$ samples to avoid major contributions from static SDW order or spin-glass-like magnetism. Given that the orbital-like magnetic order observed by spin-polarized neutron diffraction in La_{1.915}Sr_{0.085}CuO₄ is quite weak and has not been observed in other cuprates beyond p = 0.135, it is conceivable that it is not present in $x \ge 0.13$ samples. What can be said is that the ZF- μ SR measurements of La_{2-x}Sr_xCuO₄ in the Sr concentration (hole-doping) range $0.13 \le x < 0.19$ do not support theoretically predicted loop-current phases, and hence favor an alternative explanation for the unusual magnetic order detected by spin-polarized neutron diffraction at lower hole doping.

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