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Spin-orbit excitons in CoO

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CoO has an odd number of electrons in its unit cell, and therefore is expected to be metallic. Yet, CoO is strongly insulating owing to significant electronic correlations, thus classifying it as a Mott insulator. We investigate the magnetic fluctuations in CoO using neutron spectroscopy. The strong and spatially far-reaching exchange constants reported recently [P. M. Sarte et al., Phys. Rev. B 98, 024415 (2018)], combined with the single-ion spin-orbit coupling of similar magnitude [R. A. Cowley et al., Phys. Rev. B 88, 205117 (2013)] results in significant mixing between ground spin-orbit levels in the low-temperature magnetically ordered phase. The high degree of entanglement, combined with the structural domains originating from the Jahn-Teller structural distortion at ~300 K, make the magnetic excitation spectrum highly structured in both energy and momentum. We extend previous theoretical work on PrTl3 [W. J. L. Buyers et al., Phys. Rev. B 11, 266 (1975)] to construct a mean-field and multilevel spin-orbit exciton model employing the aforementioned spin exchange and spin-orbit coupling parameters for coupled Co2+ ions lying on a rocksalt lattice. This parametrization, based on a tetragonally distorted type-II antiferromagnetic unit cell, captures both the sharp low-energy excitations at the magnetic zone center, and the energy broadened peaks at the zone boundary. However, the model fails to describe the momentum dependence of the excitations at high-energy transfers, where the neutron response decays faster with momentum than the Co2+ form factor. We discuss such a failure in terms of a possible breakdown of localized spin-orbit excitons at high-energy transfers.

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I. INTRODUCTION

Mott insulators are materials where conventional band theory fails, predicting metallic behavior owing to half-filled bands, with the origin of the insulating response indicative of strong electronic correlations [1–5]. Mott insulators are parent materials for high-temperature cuprate superconductivity [6–8]. Moreover, there have been some suggestions that these insulators may even be implicated as being parent to some iron-based superconductors [9–13]. These Mott insulators display well-defined spin excitations, however, rapidly break down [14–17] on charge doping toward superconductivity [18]. More recently, Mott insulators with strong spin-orbit coupling have been of particular interest in the search for unconventional topological states [19–21]. These studies have focused on 4d and 5d transition metals with strong spin-orbit coupling resulting in j_eff = ½ ground states, and new Kitaev bond directional phases [22–24]. However, much of the single-ion physics that results in these j_eff = ½ ground states is present in Co2+-based compounds that also display strong spin-orbit coupling [25,26].

In this context, it is timely to investigate the classic Mott insulator CoO, where significant spin-orbit coupling is present and comparable to the magnetic exchange. In this study, we investigate the mixed spin-orbit transitions in CoO through their parametrization with a multilevel spin-orbit exciton model extending previous theoretical work on PrTl3 [27]. While this model reproduces the experimental data at low-energy transfers, we show that its failure at high-energy transfers is accompanied by a possible breakdown of these excitations.

For the past several decades, CoO has been one of the most extensively studied Mott insulators. The 3d metal monoxide was among the first orbitally ordered materials to be investigated with neutron diffraction [28]. Its primitive unit cell consists of one 3d7 Co2+ and one 2s2 2p6 O2−, corresponding to 15 valence electrons. With an odd number of valence electrons, conventional band theory [29,30] would predict CoO to be metallic. However, CoO is a very strong insulator [31] with a room-temperature resistivity of 108 Ω cm, and an optical band gap of 2.5 eV [32–34], with evidence for metallic behavior being found only under extremely high pressures on the order of 100 GPa [35].
distortion and various far-reaching large exchange interactions, yields a complex magnetic excitation spectrum that results from the strong entanglement of multiple spin-orbit levels [Fig. 1(c)]. The resulting multiparameter spin-orbital Hamiltonian incorporating both exchange and spin-orbit coupling of similar magnitude, further complicated by the complex magnetic ordering and structural distortions, has made the understanding of the magnetic excitations in this material particularly difficult [53].

By employing both the spin-orbit coupling constant \(\lambda\) and the magnetic exchange constants \(J\) that were experimentally determined in our previous work [54,55] on the magnetically diluted monoxide \(\text{Mg}_{0.97}\text{Co}_{0.03}\text{O}\), we will show that the low-energy magnetic excitation spectrum of CoO measured in the Néel regime by inelastic neutron spectroscopy is reproduced by a mean-field multilevel spin-orbit exciton model based on Green’s functions [27]. Our parametrization successfully captures the fine structure of the well-defined low-energy spin excitations present at the magnetic zone center and also the broadening in both momentum and energy at the zone boundaries. In contrast, the model fails to reproduce the high-energy response consisting of high-velocity excitations that decay with momentum faster than the \(\text{Co}^{2+}\) form factor. We suggest that this failure of the model provides evidence for a breakdown of localized spin-orbit excitations at high-energy transfers, possibly replaced by delocalized or itinerantlike fluctuations, despite CoO being a strong Mott insulator.

This paper is divided into three general sections. In the first section, we describe the theoretical framework that we apply to parametrize the neutron scattering response in CoO. We first outline the single-ion response defining the crystal-field Hamiltonian, and then discuss the coupled equations of motion that were used to numerically derive the neutron response. In the second section, we first present the experimental data as measured with neutron spectroscopy, followed by a direct comparison to our multilevel spin-orbit exciton model. To conclude, we discuss the high-energy excitations and the poor agreement with the \(\text{Co}^{2+}\) form factor and speculate as to their origin.

II. THEORY

We first discuss the theoretical framework used to describe the localized magnetic response in CoO. The neutron magnetic cross section is proportional to the magnetic dynamic structure factor \(S(Q, \omega)\) defined by

\[
S(Q, \omega) = g_L^2 f^2(Q) \sum_{\alpha\beta} (\delta_{\alpha\beta} - \hat{Q}_\alpha \hat{Q}_\beta) S^{\alpha\beta}(Q, \omega),
\]

where \(g_L\) is the Landé \(g\) factor, \(f(Q)\) is the magnetic form factor [50,56], and \(S^{\alpha\beta}(Q, \omega)\) corresponds to the dynamic spin structure factor. Since the orbital angular momentum is quenched (i.e., \(\langle \hat{L} \rangle = 0\)) for \(3d^7\) \(\text{Co}^{2+}\), the orbital contribution to the scattering cross section is assumed to be weak, and therefore the spin operators provide the dominant contribution to the neutron scattering cross section [57]. This assumption allows \(S^{\alpha\beta}(Q, \omega)\) to be defined in terms of expectation values of spin operators \(\hat{S}^\alpha(i, \tau)\) of index \(\nu = +, -, \text{ or } z\), acting on a site \(i\) at a time \(\tau\). Such a definition of the dynamic structure

FIG. 1. (a) First four coordination shells of the high-temperature \(\text{CoO}\) rocksalt structure. (b) Tanabe-Sugano diagram for \(3d^7\) \(\text{Co}^{2+}\) in octahedral coordination calculated by Cowley et al. [54]. Shaded rectangles correspond to experimentally measured excitations for cubic \(\text{CoO}\) at room temperature with heights and the width corresponding to experimental errors in energy and the statistical error. The dashed red line at 10\(DQ/J(=dd)\) ~ 2.5 denotes the spin crossover from (left) high-spin \(S = \frac{3}{2}\) to (right) low-spin \(S = \frac{1}{2}\). (c) Calculated normalized energy variation as a function of the tetragonal distortion \(\Delta_{\text{dis}}\) and the magnetic molecular field \(\langle \hat{H}_{\text{MF}} \rangle\) perturbations to the \(j_{\text{eff}}\) manifolds from the ground-state crystal-field triplet \(2T_1\) of \(\text{Co}^{2+}\) in octahedral coordination. Both the energy eigenvalues and individual parameters are presented to scale.

Possessing a cubic \(Fm\bar{3}m\) structure [36–40] at room temperature [Fig. 1(a)], \(\text{CoO}\) assumes long-range antiferromagnetic order at \(T_N \sim 290\) K [41], in contrast to the long-range ferromagnetism predicted by general band coupling models that assume a dominant direct exchange [42,43]. Despite being the subject of many neutron diffraction studies, its magnetic structure has proven to be particularly contentious, with both collinear [44,45] and noncollinear [46,47] models describing diffraction patterns equally well [48–52].

As illustrated in Fig. 1(b), the \(2T_1\) crystal-field ground state for \(\text{CoO}\) corresponds to the \(3d^7\) \(\text{Co}^{2+}\) assuming a high-spin \(S = \frac{3}{2}\) configuration, yielding an orbital triplet with one hole in the \(t_{2g}\) orbital manifold. The resulting orbital degeneracy, coupled with both a Jahn-Teller driven [36] unit-cell
factor is given by

$$S^{\alpha\beta}(Q, \omega) = \frac{1}{2\pi} \int dt \, e^{i\omega t} \langle \hat{S}^{\alpha}(Q, t) \hat{S}^{\beta}(-Q, 0) \rangle,$$

whose relationship to the response function $G^{\alpha\beta}(Q, \omega)$ is given by the fluctuation-dissipation theorem as

$$S^{\alpha\beta}(Q, \omega) = \frac{1}{\pi} \frac{1}{1 - \exp(\omega/k_B T)} G^{\alpha\beta}(Q, \omega). \quad (2)$$

Motivated by previous work on PrTl$_3$ [27,58], the theoretical portion of this paper begins by first writing the equations of motion for the response function in terms of commutators involving the magnetic Hamiltonian $\hat{H}$. We then investigate this magnetic Hamiltonian in CoO to define both the single-ion states and how these states are coupled from site to site on the rocksalt lattice. Finally, we apply mean-field theory to decouple the equations of motion, thereby reducing the formula for the response functions to a set of coupled linear equations that can be computed numerically and directly compared with experiment. In this approach, we use creation and annihilation operators of the single-ion states rather than the Holstein-Primakoff transformation for a single spin operator. This approach allows for both the incorporation of spin-orbit level mixing, and the explicit inclusion of the single-ion terms in the Hamiltonian, such as spin-orbit coupling, rather than employing anisotropy terms that incorporate the orbital physics through perturbation theory [59].

### A. Equation of motion for the response function

According to linear response theory, the response function measured with neutrons is proportional to the Fourier transform of the retarded Green’s function that is given by

$$G^{\alpha\beta}(i, j, t) = G(\hat{S}^{\alpha}(i, t), \hat{S}^{\beta}(j, 0)) = -i\Theta(t)\langle [\hat{S}^{\alpha}(i, t), [\hat{S}^{\beta}(j, 0), \hat{H}]] \rangle \quad \tag{3}$$

where $\Theta(t)$ is the Heaviside function. As shown in Sec. I of the Supplemental Material [60], by taking the first time derivative of $G^{\alpha\beta}(i, j, t)$, applying the Heisenberg equation of motion, and Fourier transforming from the time to the frequency domain, one arrives at the following equation of motion:

$$\omega G(\hat{A}, \hat{B}, \omega) = \langle [\hat{A}, \hat{B}] \rangle + G(\hat{A}, \hat{H}), \quad \tag{4}$$

where $\hat{A}$ and $\hat{B}$ denote generic spin operators. Equation (4) indicates that deriving a model for the neutron scattering response functions relies both on the understanding of the Hamiltonian $\hat{H}$ and its commutator with the spin operators. We now investigate the individual contributions to magnetic Hamiltonian in CoO.

### B. Total magnetic Hamiltonian $\hat{H}$

The total magnetic Hamiltonian consisting of crystal-field (CF) contributions and coupling between Co$^{3+}$ on sites $i$ and $j$ can be written as

$$\hat{H} = \hat{H}_{\text{CF}} + \sum_{ij} J(ij) \hat{S}(i) \cdot \hat{S}(j). \quad $$

By defining a molecular field Hamiltonian

$$\hat{H}_{\text{MF}}(i) = \sum_i H_{\text{MF}}(i) \hat{S}_z(i),$$

where

$$H_{\text{MF}}(i) = 2 \sum_{ij} J(ij)(\hat{S}(j) - 2\langle \hat{S}_z(j) \rangle), \quad \tag{5}$$

$\hat{H}$ can be written as a sum of a single-ion ($\hat{H}_1$) and an interion ($\hat{H}_2$) term given by

$$\hat{H}_1 = \sum_i \hat{H}_{\text{CF}}(i) + \sum_i \hat{S}_z(i) \left( 2 \sum_{ij} J(ij)(\hat{S}(j)) \right) \quad \tag{6}$$

and

$$\hat{H}_2 = \sum_{ij} J(ij)(\hat{S}_z(i)\hat{S}_z(j) - 2\langle \hat{S}_z(j) \rangle)$$

$$+ \frac{1}{2} \sum_{ij} J(ij)[\hat{S}_+(i)\hat{S}_-(j) + \hat{S}_-(i)\hat{S}_+(j)], \quad \tag{7}$$

where $\langle \hat{S}_z(j) \rangle$ denotes a thermal average given by

$$\langle \hat{S}_z(n) \rangle = \sum_n f_n \langle n | \hat{S}_z | n \rangle \equiv \sum_n \hat{S}_{azon} f_n, \quad \tag{8}$$

with $\hat{S}_{azon} = \langle n | \hat{S}_z | n \rangle$, and $f_n$ is the Boltzmann thermal population factor. The inclusion of a factor of 2 in Eq. (6) follows the convention that was established in Ref. [53] to explicitly account for the double counting in the sum over sites.

The procedure we follow to derive the neutron response consists of two parts. First, we diagonalize the single-ion component $\hat{H}_1$ for a given molecular field such that

$$\hat{H}_1 = \sum_i \sum_n \omega_n C_n^i(i) C_n(i), \quad \tag{9}$$

where $C(i)$ and $C'(i)$ are ladder operators satisfying the commutation relations $[C_n(i), C_{n'}(i')] = \delta_{n,n'} \delta_{i,i'}$, and $\omega_n$ are the energy eigenvalues. The second step consists of using these states to apply mean-field theory on the interion $\hat{H}_2$ term to then compute the neutron response using the equation of motion given in Eq. (4). We will now discuss the eigenstates of the single-ion component of the Hamiltonian $\hat{H}$, followed by the interion component. This section concludes with the application of mean-field theory on the interion component, allowing for the derivation of an expression for the Green’s function that can be calculated numerically and compared directly to experiment.

### 1. Single-ion Hamiltonian $\hat{H}_1$

As schematically illustrated in Fig. 1(c), the single-ion component of the Hamiltonian consists of four components

$$\hat{H}_1 = \hat{H}_{\text{CF}} + \hat{H}_{\text{MF}}$$

$$= (\hat{H}_{\text{CEF}} + \hat{H}_{\text{SO}} + \hat{H}_{\text{dis}}) + \hat{H}_{\text{MF}},$$

corresponding to the contributions from the crystalline electric field $\hat{H}_{\text{CEF}}$, spin-orbit $\hat{H}_{\text{SO}}$, structural distortion $\hat{H}_{\text{dis}}$, and mean molecular field $\hat{H}_{\text{MF}}$. Hyperfine nuclear transitions are neglected since previous measurements [61] have indicated
that these are on the order of $\sim \mu$eV, and thus beyond the experimental resolution of our study. We now discuss each term of the Hamiltonian $\hat{\mathcal{H}}_1$.

Crystalline electric field, $\hat{\mathcal{H}}_{\text{CEF}}$. As illustrated in the Tanabe-Sugano [62–64] diagram in Fig. 1(b), for the case of CoO, the $3d^7 \text{Co}^{3+}$ is octahedrally coordinated by the weak field $\text{O}^{2-}$ ligand resulting in a crystal-field splitting $10Dq$ that is weaker than the energy differences between the free-ion terms [65]. Consequently, the crystalline electric field contribution $\hat{\mathcal{H}}_{\text{CEF}}$ can be treated as a perturbation to the free-ion basis states that are defined by Hund’s rules incorporating the effects of electron-electron Coulomb repulsion and the Pauli exclusion principle. A combination of Hund’s first rule of maximum multiplicity and second rule requiring the total orbital angular momentum $L$ be maximized yields a total spin of $S = \frac{5}{2}$ and orbital angular momentum $L = 3$, for the $3d^7 \text{Co}^{3+}$, corresponding to an orbital ground-state term symbol of $^4F$.

The crystalline electric field $\hat{\mathcal{H}}_{\text{CEF}}$ contribution corresponding to the octahedral coordination of the $^4F$ free-ion ground state by $\text{O}^{2-}$ ligands can be expressed in terms of the Stevens operators $\hat{\mathcal{O}}^4_0$ and $\hat{\mathcal{O}}^4_4$, and the numerical coefficient $B_d < 0$ [66,67] as

$$\hat{\mathcal{H}}_{\text{CEF}} = B_d (\hat{\mathcal{O}}^4_0 + 5\hat{\mathcal{O}}^4_4).$$

Since the spin-orbit coupling is expected to be considerably weaker than the crystal-field contribution for the $3d$ Co$^{3+}$, the complete set of commuting observables are $L^2$, $L_z$, $S^2$, and $\hat{\mathcal{S}}$, with corresponding good quantum numbers $L$, $m_L$, $S$, and $m_S$ in the Russell-Saunders $L$-$S$ coupling scheme; thus, by the Wigner-Eckart theorem, both Stevens operators in $\hat{\mathcal{H}}_{\text{CEF}}$ can be defined in the $|L, m_L\rangle$ basis, as summarized in Sec. II of the Supplemental Material [60].

The diagonalization of $\hat{\mathcal{H}}_{\text{CEF}}$ results in an orbital triplet ground state ($^4T_1$), an excited orbital triplet ($^4T_2$), and an orbital singlet $^4A_2$, where $\Delta(^4T_1 \rightarrow ^4T_2) = 480B_d$ and $\Delta(^4T_2 \rightarrow ^4A_2) = 600B_d$. The Stevens factor $B_d$ is related to the crystal-field splitting by $10Dq = 400B_d$ [Fig. 1(b)], where $10Dq$ was previously measured to be $\sim 1$ eV [54,68–72]. Since the $^4T_1$ crystal-field ground state and $^4T_2$ first excited state are separated by $\sim$eV, it is a valid approximation that the $^4T_1$ ground state will exclusively determine the magnetic properties of CoO [73,74].

Spin-orbit coupling, $\hat{\mathcal{H}}_{\text{SO}}$. The second perturbation to the $^4F$ free-ion ground state is spin-orbit coupling given by

$$\hat{\mathcal{H}}_{\text{SO}} = \lambda \hat{\mathcal{L}} \cdot \hat{\mathcal{S}},$$

where $\lambda$ is the spin-orbit coupling constant. A common approach is to exclusively consider the $^4T_1$ ground state, requiring a projection from the original $|L = 3, m_L\rangle$ basis onto a smaller basis $|l = 1, m_l\rangle$ that defines the subspace that is spanned by the crystal-field ground state. As discussed by Abragam and Bleaney [66], this particular projection can be performed using representation theory. Here, we outline an alternate method based on the matrix representation of angular momentum operators [75] that was inspired by the work on $4d$ and $5d$ transition metal oxides by Stamokostas and Fiete [76].

The matrix approach begins by first determining the set of eigenvectors $|\phi_{\text{CEF}}\rangle$ of the crystalline electric field Hamiltonian $\hat{\mathcal{H}}_{\text{CEF}}$ in the $|L = 3, m_L\rangle$ basis. Since $|\phi_{\text{CEF}}\rangle$ is also a basis, a transformation matrix $C$ can be constructed that rotates from the $|L = 3, m_L\rangle$ to the $|\phi_{\text{CEF}}\rangle$ basis. The matrix $C$ consists of columns corresponding to eigenvectors of $\hat{\mathcal{H}}_{\text{CEF}}$ in the $|L = 3, m_L\rangle$ basis arranged in order of increasing energy eigenvalues. In the case of degenerate eigenvalues, a small perturbative Zeeman term of the form $\epsilon \hat{\mathcal{S}}_z$, with $\epsilon$ being a small constant, was applied to remove the degeneracy and uniquely define the column order. For Co$^{3+}$ in octahedral coordination with $B_d$ set to $-1$, $C$ is given by

$$C = \begin{bmatrix}
0 & 0 & -0.79 & 0.61 & 0 & 0 & 0 \\
0 & 0 & 0 & 0 & -0.71 & 0 & -0.71 \\
0.61 & 0 & 0 & 0 & 0 & -0.79 & 0 \\
0 & 0 & 0 & 0 & 0 & 0 & 0 \\
1.00 & 0 & 0 & 0 & 0 & 0 & 0 \\
0 & 0 & -0.61 & -0.79 & 0 & 0 & 0 \\
0 & 0 & 0 & 0 & -0.71 & 0 & 0.71 \\
0 & 0 & 0 & 0 & 0 & 0 & 0.61 & 0
\end{bmatrix}.$$
confirming a doublet ground state with quartet and sextet excited manifolds [Fig. 1(c)] with
the subspaces spanned by the spin-orbit manifolds of $\hat{L}_z$.

By assigning an effective angular momentum operator to the subspace spanned by the $|l = 1, m_l\rangle$ basis, it is implied these new operators must follow the same commutation relations for general angular momentum operators. To check this fundamental requirement is satisfied, we have transformed the three $\hat{L}_x, \hat{L}_y, \hat{L}_z$ operators, each of which is a $7 \times 7$ matrix, to the $\phi_{\text{CEF}}$ basis with the procedure outlined above. We then extracted the top $3 \times 3$ block matrices of the projected matrices $C^{-1}L_{x,y,z}C$ to define $\hat{L}_{x,y,z}$, and confirmed that these matrices do follow the commutation relations of angular momentum given by $A \times A = i$. We note that the presence of a thermally isolated low-energy triplet does not guarantee that these commutation relations are followed. An example of such a failure has been recently discussed in the context of low-energy doublets in the heavy fermion CeRhSi$_3$ [78].

Having projected $\hat{L}$ onto a fictitious operator $\hat{I}$ to reflect the triplet orbital degeneracy of the $^4T_1$ ground state of $\hat{H}_{\text{CEF}}$, we now derive the eigenstates of the perturbative $\hat{H}_{\text{SO}}$ term. The basis is now the $12 \times 12$ matrix formed the three $\hat{L}_x, \hat{L}_y, \hat{L}_z$ operators, each of which is a $7 \times 7$ matrix, to the $\phi_{\text{CEF}}$ basis with the procedure outlined above. We then extracted the top $3 \times 3$ block matrices of the projected matrices $C^{-1}L_{x,y,z}C$ to define $\hat{L}_{x,y,z}$, and confirmed that these matrices do follow the commutation relations of angular momentum given by $A \times A = i$. We note that the presence of a thermally isolated low-energy triplet does not guarantee that these commutation relations are followed. An example of such a failure has been recently discussed in the context of low-energy doublets in the heavy fermion CeRhSi$_3$ [78].

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A comparison of the block matrices of the projected $\hat{J}_2$ operator as given above to the $\hat{J}_1$ operator in the $|\vec{J}_1\rangle = |\frac{1}{2}, \frac{1}{2}, \frac{1}{2}\rangle$, $|\vec{J}_2\rangle = |\frac{1}{2}, \frac{1}{2}, \frac{1}{2}\rangle$, and $|\vec{J}_3\rangle = |\frac{3}{2}, \frac{3}{2}, \frac{3}{2}\rangle$ bases confirms that the top 2 x 2, middle 4 x 4, and bottom 6 x 6 block matrices correspond to $|\vec{J}_1\rangle = |\frac{1}{2}, \frac{1}{2}, \frac{1}{2}\rangle$, and $|\vec{J}_3\rangle = |\frac{3}{2}, \frac{3}{2}, \frac{3}{2}\rangle$, respectively. By projecting the $\hat{J}_1$ and $\hat{J}_2$ operators, it can be shown that these block matrices also satisfy the canonical commutation relations of angular momentum.

**Distortion Hamiltonian, $\hat{H}_\text{dis}$**. The next perturbation to the single-ion Hamiltonian corresponds to the structural deformation of the CoO unit cell that accompanies long-range antiferromagnetic order, resulting in the distortion of the crystalline electric field from ideal octahedral coordination [79]. While the exact symmetry of the low-temperature phase has proven to be particularly contentious, we will consider a simple tetragonal distortion, corresponding to a uniaxial distortion parameter $\alpha$. In these particular magnets, the degree of mixing is much weaker. In these particular magnets, the degree of mixing is much weaker. In these particular magnets, the degree of mixing is much weaker. In these particular magnets, the degree of mixing is much weaker. In these particular magnets, the degree of mixing is much weaker. In these particular magnets, the degree of mixing is much weaker. In these particular magnets, the degree of mixing is much weaker. 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FIG. 2. (a) Background (pure and nonmagnetic MgO) subtracted powder-averaged neutron-scattering intensity maps of Mg$_{0.97}$Co$_{0.03}$O measured on (top) MARI at 5 K with an $E_i$ = 30 meV, (middle) MARI at 5 K with an $E_i$ = 10 meV, and (bottom) IRIS at 11 K with an $E_f$ of 1.84 meV revealing seven low-energy bands of dispersionless magnetic excitations. (b) Relevant energy scales for the effective pair Hamiltonian. (c) Calculated difference in the ground-state manifold’s energy eigenstates obtained from the diagonalization of the effective pair Hamiltonian (black line). The nonlinearity is in contrast with the behavior predicted by the projection theorem (gray line). (Inset) The mechanism for antiferromagnetism (top) and weaker ferromagnetism (bottom) is a result of a combination of the 90° Co$^{2+}$-O$^{2-}$-Co$^{2+}$ exchange pathway and the orbital degree of freedom in the $t_{2g}$ channel on each Co$^{2+}$, in agreement with the predictions of the Goodenough-Kanamori-Anderson rules. Yellow arrows denote local $t_{2g}$ spin configurations and teal arrows denote total spin configurations on each Co$^{2+}$.

rotation being given by

$$S_{(±,z)} = \sum_{mn} S_{(±,z)mn} C_m^\dagger C_n.$$  \hspace{1cm} (12)

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Initial value</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\lambda$</td>
<td>$-16$</td>
<td>[54]</td>
</tr>
<tr>
<td>$\Gamma$</td>
<td>$-8.76$</td>
<td>[101,102]</td>
</tr>
<tr>
<td>$J_{1F}$</td>
<td>$-0.918$</td>
<td>[55]</td>
</tr>
<tr>
<td>$J_{1AF}$</td>
<td>$1.000$</td>
<td></td>
</tr>
<tr>
<td>$J_2$</td>
<td>$3.09$</td>
<td></td>
</tr>
<tr>
<td>$J_3$</td>
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<td></td>
</tr>
<tr>
<td>$J_{3AF}$</td>
<td>$0.262$</td>
<td></td>
</tr>
<tr>
<td>$J_{4F}$</td>
<td>$-0.0504$</td>
<td></td>
</tr>
<tr>
<td>$J_{4AF}$</td>
<td>$0.0759$</td>
<td></td>
</tr>
<tr>
<td>$H_{MF}$</td>
<td>$64.8$</td>
<td>[98–100]</td>
</tr>
</tbody>
</table>

TABLE I. Initial values (in meV) for the parameters of the spin-orbit exciton model.

By writing the full Hamiltonian $\hat{H} = \hat{H}_1 + \hat{H}_2$ in terms of the ladder operators as defined in Eq. (12), and using the definition of the interlevel susceptibility $G$ defined by

$$G^{\alpha\beta}(i, j, \omega) = \sum_{mn} S_{(±,z)mn} G^{\beta}(m, n, i, j, \omega),$$  \hspace{1cm} (13)

where the indices $\alpha, \beta$ are either $+, -, or z$, the second term on the right-hand side of the equation of motion of the Green’s function [Eq. (4)] reduces to three sets of commutators, termed diagonal, transverse, and longitudinal, with each involving spin operators rewritten in terms of ladder operators, as discussed in Sec. III of the Supplemental Material [60]. Buyers et al. [27] demonstrated that by combining the random phase decoupling method [90–93] (Sec. IV of the Supplemental Material [60]) with the definitions of both the single-site response function

$$g^{\alpha\beta}(E) = \sum_{n} \left\{ \frac{S_{\alpha\beta}}{E_i + \Delta - E_{n0} - \frac{S_{\alpha\beta}}{E_i + \Delta + E_{n0}}} \right\},$$  \hspace{1cm} (14)
and the Fourier transform of the exchange interaction \( J(\mathbf{Q}) \)
\[
J(\mathbf{Q}) = \sum_{i \neq j} J_{ij} e^{i \mathbf{Q} \cdot \mathbf{d}_{ij}},
\]
(15)

the full neutron response Fourier transformed into momentum \( \mathbf{Q} \) space can be written as a set of coupled linear equations given by
\[
G^{\alpha \beta}(\mathbf{Q}, E) = e^{i \mathbf{Q} \cdot \mathbf{d}}(E) + e^{i \mathbf{Q} \cdot \mathbf{d}}(E)J(\mathbf{Q})G^{-\beta}(\mathbf{Q}, E)
+ e^{i \mathbf{Q} \cdot \mathbf{d}}(E)J(\mathbf{Q})G^{+\beta}(\mathbf{Q}, \omega)
+ 2 e^{i \mathbf{Q} \cdot \mathbf{d}}(E)J(\mathbf{Q})G^{\beta\beta}(\mathbf{Q}, E),
\]
(16)

where \( \omega \) has been relabeled as \( E = \hbar \omega \). Here, we have employed the \( T \to 0 \) K single-site response function since the energy transfers under consideration in the current investigation (\( \geq 20 \) meV) are much larger than the sample temperature (\( \sim 0.5 \) meV). The denominator of \( g^{\alpha \beta} \) consists of \( E_{mm} = \hbar \omega_m - \hbar \omega_0 \) corresponding to the energy associated with a transition from the ground state \( |0\rangle \) to the \( |n\rangle \) eigenstate of the single-ion Hamiltonian \( \mathcal{H}_1 \), while the presence of the positive infinitesimal \( \Delta \) is to ensure analyticity, and was set to 50% of the experimental resolution width (HWHM) on MERLIN [94] that was calculated by PYCHOP [95]. Coupling between the single-site response functions, and thus the dispersion of the total response functions \( G^{\alpha \beta} \), is defined by \( J(\mathbf{Q}) \) which is parametrized by both \( J_{ij} \) and \( \mathbf{d}_{ij} \), denoting the exchange constant and displacement vector, respectively, between sites \( i \) and \( j \). As a first approximation, our calculations have considered the simplest case where the exchange interaction is spatially isotropic. We note that in general this is not the case, owing to the anisotropy of the orbital configuration of Co\(^{2+}\).

By considering all possible combinations of indices \( \alpha, \beta \) in Eq. (16) and noting that the nonzero single-site response functions for Co\(^{2+}\) in such a highly symmetric environment are \( g^{++}, g^{-+}, \) and \( g^{--} \), only three nonzero Green’s functions are obtained:
\[
G^{++}(\mathbf{Q}, E) = g^{++}(E) + g^{++}(E)J(\mathbf{Q})G^{++}(\mathbf{Q}, E),
G^{-+}(\mathbf{Q}, E) = g^{-+}(E) + g^{-+}(E)J(\mathbf{Q})G^{-+}(\mathbf{Q}, E),
G^{--}(\mathbf{Q}, E) = g^{--}(E) + 2g^{--}(E)J(\mathbf{Q})G^{--}(\mathbf{Q}, E),
\]
(17)

with both \( G^{++} \) and \( G^{--} \) being both zero, as required by definition of the retarded Green’s function.

The simplest model for the long-range antiferromagnetic order in CoO is a type-II collinear antiferromagnetic magnetic structure [28]. Corresponding to (111) ferromagnetic sheets stacked antiferromagnetically along the [111] direction, this type of magnetic structure has been observed in CoO under pressure, despite the suppression of the structural distortion [96]. Such a model implies that CoO can be reduced to two unique magnetic sublattices; thus, the site indices \( i \) and \( j \) assume labels of either 1 or 2, and Eq. (17) becomes four coupled linear equations
\[
G_{11}^{++}(\mathbf{Q}, E) = g_{11}^{++}(E) + g_{11}^{++}(E)J_1(\mathbf{Q})G_{11}^{++}(\mathbf{Q}, E)
+ g_{11}^{++}(E)J_2(\mathbf{Q})G_{21}^{++}(\mathbf{Q}, E),
G_{21}^{++}(\mathbf{Q}, E) = g_{21}^{++}(E)J_1(\mathbf{Q})G_{12}^{++}(\mathbf{Q}, E)
+ g_{21}^{++}(E)J_2(\mathbf{Q})G_{22}^{++}(\mathbf{Q}, E),
G_{12}^{++}(\mathbf{Q}, E) = g_{12}^{++}(E)J_1(\mathbf{Q})G_{12}^{++}(\mathbf{Q}, E)
+ g_{12}^{++}(E)J_2(\mathbf{Q})G_{22}^{++}(\mathbf{Q}, E),
G_{22}^{++}(\mathbf{Q}, E) = g_{22}^{++}(E) + g_{22}^{++}(E)J_1(\mathbf{Q})G_{12}^{++}(\mathbf{Q}, E)
+ g_{22}^{++}(E)J_2(\mathbf{Q})G_{22}^{++}(\mathbf{Q}, E),
\]
(18)

and
\[
G_{11}^{--}(\mathbf{Q}, E) = g_{11}^{--}(E) + 2g_{11}^{--}(E)J_1(\mathbf{Q})G_{11}^{--}(\mathbf{Q}, E)
+ 2g_{11}^{--}(E)J_2(\mathbf{Q})G_{21}^{--}(\mathbf{Q}, E),
G_{21}^{--}(\mathbf{Q}, E) = 2g_{21}^{--}(E)J_1(\mathbf{Q})G_{12}^{--}(\mathbf{Q}, E)
+ 2g_{21}^{--}(E)J_2(\mathbf{Q})G_{22}^{--}(\mathbf{Q}, E),
G_{12}^{--}(\mathbf{Q}, E) = 2g_{12}^{--}(E)J_1(\mathbf{Q})G_{12}^{--}(\mathbf{Q}, E)
+ 2g_{12}^{--}(E)J_2(\mathbf{Q})G_{22}^{--}(\mathbf{Q}, E),
G_{22}^{--}(\mathbf{Q}, E) = g_{22}^{--}(E) + 2g_{22}^{--}(E)J_1(\mathbf{Q})G_{12}^{--}(\mathbf{Q}, E)
+ 2g_{22}^{--}(E)J_2(\mathbf{Q})G_{22}^{--}(\mathbf{Q}, E),
\]
(19)

with \( J_1 \) and \( J_2 \) denoting \( J(\mathbf{Q}) \) on the same \( i = j \) and different \( i \neq j \) sublattices, respectively. Solving these four coupled equations yields

\[
G^{++}(\mathbf{Q}, E) = \sum_{ij} G_{ij}^{++}(\mathbf{Q}, E) = \frac{g_{11}^{++}(E) + g_{22}^{++}(E) + 2g_{12}^{++}(E)g_{21}^{++}(E)}{1 - g_{11}^{++}(E)J_1(\mathbf{Q})} + \frac{g_{12}^{++}(E)}{1 - g_{22}^{++}(E)J_2(\mathbf{Q})}J_2(\mathbf{Q}),
\]

\[
G^{--}(\mathbf{Q}, E) = \sum_{ij} G_{ij}^{--}(\mathbf{Q}, E) = \frac{g_{11}^{--}(E) + g_{22}^{--}(E) + 4g_{12}^{--}(E)g_{21}^{--}(E)}{1 - g_{11}^{--}(E)J_1(\mathbf{Q})} + \frac{4g_{12}^{--}(E)}{1 - g_{22}^{--}(E)J_2(\mathbf{Q})}J_2(\mathbf{Q}),
\]

where \( G^{++}(\mathbf{Q}, E) \) has the same form as \( G^{++}(\mathbf{Q}, E) \) with indices \( + \leftrightarrow - \). The equations above are a function of the single-site response function \( g^{\alpha \beta} \) and the Fourier transform of the exchange interaction \( J(\mathbf{Q}) \).

In contrast to the single-site response function, \( J(\mathbf{Q}) \) does explicitly depend on the particular magnetic sublattice under consideration, stemming from the presence of the position indices \( i \) and \( j \) in its definition given by Eq. (15). The Co\(^{2+}\) sites in the \( d \) and \( s \) sublattices for a particular coordination shell \( m \) was determined by first selecting a reference Co\(^{2+}\) cation, thus defining a reference (111) plane as illustrated in Fig. 3. By definition of the type-II antiferromagnetic structure, Co\(^{2+}\) cations located on odd integer number of (111) planes away from the reference plane are defined as belonging to the \( d \) sublattice, while Co\(^{2+}\) cations located in the same or an even integer number of (111) planes away are on the \( s \) sublattice.
The program VESTA [97] was used to determine the displacement vectors to calculate $J_{s,d}$ from Eq. (15) giving

$$ J_{s}(Q) = 2J_1[\cos(\pi (H-K)) + \cos(\pi (K-L)) + \cos(\pi (L-H))] + \cdots \quad (m = 1) $$
$$ 2J_2[\cos(2\pi H) + \cos(2\pi K) + \cos(2\pi L)] + \cdots \quad (m = 2) $$
$$ 2J_3[\cos(\pi (2H-K-L)) + \cos(\pi (2H-K-L)) + \cos(\pi (2H+K+L))] + \cdots \quad (m = 3) $$
$$ \cos(\pi (H+L)) + \cos(\pi (H+K)) + \cos(\pi (K+L))] \quad (m = 4) $$

and

$$ J_{d}(Q) = 2J_1[\cos(\pi (H+K)) + \cos(\pi (K+L)) + \cos(\pi (L+H))] + \cdots \quad (m = 1) $$
$$ 2J_2[\cos(2\pi H) + \cos(2\pi K) + \cos(2\pi L)] + \cdots \quad (m = 2) $$
$$ 2J_3[\cos(\pi (2H-K+L)) + \cos(\pi (2H+K-L)) + \cos(\pi (2H-K-L))] + \cdots \quad (m = 3) $$

where the contributions from each coordination shell $m$ have been labeled explicitly.

By employing the definitions of the single-site response function $g^{\alpha\beta}$ [Eq. (14)], and the Fourier transform of the exchange interaction $J_{s,d}(Q)$ [Eqs. (20) and (21)], the total response function $G(Q, E)$ given by

$$ G(Q, E) \equiv \sum_{\alpha\beta} G^{\alpha\beta}(Q, E) $$
$$ = G^{\pm\pm}(Q, E) + G^{\pm-}(Q, E) + G^{-\pm}(Q, E) \quad (22) $$

having been labeled explicitly.

![Diagram](image)

FIG. 3. Isometric view of all Co$^{2+}$ cations located in the (a) first ($m = 1$), (b) second ($m = 2$), (c) third ($m = 3$), and (d) fourth ($m = 4$) coordination shells of the CoO rocksalt structure. For the purposes of reference, all (111) planes are labeled as either $s$ and $d$ planes with respect to the reference Co$^{2+}$ (central black site). All displacement vectors $\mathbf{a}_{\alpha\beta}$ are listed in Table SI in the Supplemental Material [60] and used to calculate $J_{s}(Q)$ and $J_{d}(Q)$ discussed in the text.

can be calculated numerically. In the $T \to 0$ K limit, the imaginary part of $G(Q, E)$ is proportional to the dynamical structure factor [Eq. (2)], and thus Eq. (1) may be reduced to

$$ S(Q, E) \propto -f^2(Q)G(Q, E), $$

demonstrating that the imaginary component of the total response function given by Eq. (22), with the inclusion of the magnetic form factor which here has been approximated by the isotropic magnetic form factor $f(Q)$, is directly proportional to the neutron magnetic cross section.

D. Parameters: Initial values and orbital configurations

Having presented our model, we now discuss the parametrization of the spin-orbit excitations in CoO. Since our model approximates CoO as a tetragonally distorted type-\text{II} antiferromagnet, the single-site response $g^{\alpha\beta}$, itself being a function of the single-ion Hamiltonian $\hat{H}_i$ [Eq. (6)], is defined by three parameters: $\lambda$, $\Gamma$, and $H_{\text{MF}}$. The spin-orbit coupling parameter $\lambda$ was taken to be $-16$ meV, corresponding to its value reported by Cowley et al. [54]. An initial estimate for the mean molecular field $H_{\text{MF}}$ was determined by first extracting the value for $\sum_{i>j} z_{ij}J_{ij}$ from the reported [98–100] Curie-Weiss temperature $\theta_{\text{CW}} = -330$ K ($-28.4$ meV) via its mean-field definition

$$ \theta_{\text{CW}} = -\frac{2}{3\xi}S(S+1)\sum_{i>j} z_{ij}J_{ij}, \quad (23) $$

where $\xi$ is a scale factor of 1.9 calculated by Kanamori [98] accounting for mixing between the $3^+$ and $4^+$ free-ion states. Inserting the value of $\sum_{i>j} z_{ij}J_{ij}$ into the definition of $H_{\text{MF}}$ given by Eq. (5) yields an initial estimate of $64.8$ meV. An initial estimate of the tetragonal distortion parameter $\Gamma = -8.76$ meV was determined by scaling the value of $-1.49$ meV that was reported for KCoF$_3$ [101] by an empirical factor of 0.0116/0.00197 = 5.89 corresponding to the ratio of their respective tetragonal distortions $\delta a/a$.

To define $J(Q)$, we have taken the values for the exchange constants for the dilute monoxide Mg$_{0.97}$Co$_{0.03}$O [55] as estimates for pure undiluted CoO since these exchange constants correspond to a Curie-Weiss temperature [Eq. (23)]
in close agreement with the value reported for CoO. However, our investigation on Mg$_{0.97}$Co$_{0.03}$O also revealed that each coordination shell possessed the possibility for both antiferromagnetic and ferromagnetic coupling, with the exception of the second nearest neighbor which is fixed to be antiferromagnetic by the 180° Co$^{2+}$-O$^{2-}$-Co$^{2+}$ superexchange pathway. As illustrated in Fig. 2(c), inset, such dual behavior is a direct consequence of the $t_{2g}$ degeneracy of the high-spin $d^7$ configuration of Co$^{2+}$, and thus a particular choice of $J$, be it antiferromagnetic or ferromagnetic, corresponds to a specific local orbital configuration. By incorporating this dual behavior for coordination shells 1, 3, and 4, we must consider all $2^3 = 8$ sets of exchange constants of the form $x\Lambda x\gamma$, where $x$ can be either antiferromagnetic (A) or ferromagnetic (F). Furthermore, since our model incorporates the effects of a tetragonal (uniaxial) distortion, we must also distinguish the involvement of a distorted or undistorted bonding configuration for each of the 8 $x\Lambda x\gamma$ orbital configurations. Thus, with these 2 additional degrees of freedom, our model must consider 16 different orbital configurations of the form $x\Lambda x\gamma\psi\phi\sigma$, where the index $\gamma = 1$ or 2, distinguishing the presence or absence of distorted bonding configurations.

Each of these 16 $x\Lambda x\gamma\psi\phi\sigma$ combinations may be interpreted as a unique orbital configuration, physically corresponding to a unique type of “domain” in the bulk CoO single crystal, each of which is subject to a different mean molecular field $H_{\text{MF}}$. In contrast, since all cations under consideration are assumed to be Co$^{2+}$ in octahedral coordination subject to a cooperative Jahn-Teller tetragonal distortion, $\lambda$, $\Gamma$, and the individual $J_n$ values (where $n$ denotes a particular type of coupling in a coordination shell $m$) are fixed to be equal for each of the 16 $x\Lambda x\gamma\psi\phi\sigma$ orbital configurations. By noting that our neutron spectroscopic measurements were performed with a large experimental beam that irradiated a macroscopic number of domains in the single crystal of CoO, our model considers the mean contribution from all 16 equally weighted $x\Lambda x\gamma\psi\phi\sigma$ orbital configurations. The initial parameters considered in the model and their initial values are summarized in Table I.

III. EXPERIMENT

Having discussed both the underlying theory of Co$^{2+}$ magnetism and the corresponding physical parameters that constitute our spin-orbit exciton model, we will now address the experimental results from neutron scattering experiments on a single crystal of CoO. This section begins with a description of the experimental techniques, followed by a summary of the neutron spectroscopic data. We conclude this section with a description of how our model was used to interpret the low-energy fluctuations of CoO deep within the Néel regime.

A. Experimental details

Sample preparation. Polycrystalline samples of CoO were synthesized by annealing high purity Co$_3$O$_4$ (>99.99%) under flowing Ar at 1200 °C for 36 h with intermittent grinding until laboratory x-ray diffraction confirmed the absence of the Co$_3$O$_4$ precursor. The phase pure CoO powder was compressed into cylindrical rods using a hydraulic press and subsequently annealed under flowing Ar at 1275 °C for 24 h in a horizontal annealing furnace. Crystal growth was performed using the floating zone technique with a four-mirror optical floating zone halogen furnace (CSI system Inc.), yielding a 10-g single crystal of CoO ($l = 50$ mm, $\phi = 8$ mm). The feed and seed rods were counter-rotated at 35 rpm with a vertical translation of 2 to 4 mm h$^{-1}$ in a pure Ar atmosphere. The initial polycrystalline seed rod was replaced for subsequent runs by single-crystal seeds from earlier growths. Previous [54] optical and scanning electron microscopy, x-ray diffraction and DC susceptibility measurements on the single crystal confirmed the presence of a single growth domain with a mosaic spread of approximately 0.1° and the absence of both magnetic or strain domains on the crystal surface and Co$_3$O$_4$ impurities, respectively.

Neutron inelastic scattering. Neutron inelastic scattering measurements were performed on the MERLIN direct geometry chopper spectrometer [94] at the ISIS neutron spallation source (Didcot, UK). The $I_0$ chopper was spun at 50 Hz in parallel with the “sloppy” Fermi chopper package to fix the incident energy with the energy transfer defined as $E = E_i - E_f$. To access a large dynamic range, three fixed incident energies $E_i$ of 110, 75, and 45 meV were selected with Fermi chopper frequencies of 350, 300, and 250 Hz, providing a resolution at the elastic line ($E = 0$) of 7.3, 4.8, and 2.7 meV, respectively. A 5-g portion of the CoO single crystal was mounted in a top-loading closed-cycle refrigerator such that the [110] and [001] crystallographic axes lay within the horizontal plane. A tomographic reconstruction in momentum space was accomplished by rotating the crystal about the [0$\bar{1}$0] axis over 120° in 0.5° steps.

The four-dimensional ($Q, E$) experimental data at each angle $\Psi$ and $E_i$ was collected at 5 K for an accumulated charge of 30 $\mu$A h on the spallation target. The raw experimental data were normalized by accumulated proton charge, corrected for detector-efficiency using a vanadium reference sample, and reduced by the MANTID data analysis software [103,104]. Visualization and manipulation of reduced experimental data including rebinning and projections were performed using the HORACE software package distributed by ISIS [105].

B. Experimental data

We begin by first presenting a summary of the experimental data from single-crystal neutron spectroscopic measurements allowing for a direct comparison with previous work on CoO to establish consistency. The experimental data from the MERLIN chopper spectrometer at 5 K is presented in Fig. 4 in the form of ($Q, E$) slices along both (1.5 ± 0.1, 1.5 ± 0.1, $L$) [Figs. 4(a) and 4(c)] and (2.0 ± 0.1, 2.0 ± 0.1, $L$) [Fig. 4(b)] capturing both the magnetic zone center and boundary, where both are compared to previously published work in the form of $Q$-integrated cuts presented in Figs. 4(d) and 4(e), respectively. As illustrated in Fig. 4(a), a ($Q, E$) slice with an incident energy $E_i = 110$ meV exhibits a band of excitations extending from ~20 meV up to ~60 meV energy transfer, corresponding to a similar range in energy transfer reported by previous THz [107] and Raman [108] spectroscopic measurements. These excitations decrease in intensity with increasing $L$, as is expected for the Co$^{2+}$ magnetic form factor, thus indicating these excitations are possibly magnetic. A higher
FIG. 4. \( (Q, E) \) slices of CoO measured on MERLIN at 5 K with an \( E_i \) of (a) 110 meV, (b) 75 meV, and (c) 45 meV. All \( (Q, E) \) slices have been folded along [001]. A comparison of \( Q \)-integrated cuts of (c) and (b) with previous measurements in the literature at the (d) magnetic zone centers, and (e) magnetic zone boundaries, respectively. Solid lines in (e) indicate the location of excitations previously determined by IR spectroscopy [106]. Horizontal bars indicate instrumental resolution.

resolution slice with an \( E_i = 45 \text{ meV} \) [Fig. 4(c)] reveals that the band of excitations corresponds to a fine structure consisting of a series of modes that are unevenly spaced in energy, in agreement with previous triple-axis [53,109], time-of-flight data [110], and Raman spectroscopy [111], with the exception of a broader peak reported for triple-axis measurements at \( \sim 40 \text{ meV} \). However, it is important to note that the previously reported triple-axis measurements employed final energies \( E_f = 14.6 \text{ and } 30.5 \text{ meV} \), both of which potentially produce spurious signals near 40 meV as a result of weak elastic scattering corresponding to \( E_i \rightarrow 4 E_f (\lambda_f/2) \) and \( 4 E_f (\lambda_f/2) \rightarrow 9 E_f (\lambda_f/3) \) [112]. This weak elastic process may have contributed to the extra scattering intensity that was observed in the previously reported triple-axis data, yet is absent in our current time-of-flight data. We also note the presence of nearly dispersionless optical phonon branches near this energy may also contribute to the overall neutron cross section [113,114]. We will later discuss how this fine structure near the magnetic zone center can be understood in terms of the spin-orbit excitations.

Slices and \( Q \)-integrated cuts through the magnetic zone boundary presented in Figs. 4(b) and 4(e) show additional complexity present in the neutron response. At low-energy transfers below \( \sim 20 \text{ meV} \), strong acoustic phonons can be seen in Fig. 4(b) to disperse from the even integer positions in momentum transfers. The phonon nature of these excitations is confirmed by the fact that the cross section grows with increasing momentum transfer \( L \), in contrast to the magnetic cross section that is subject to the Co \( 2^+ \) form factor. In addition to the acoustic phonons, a flat band is observed at \( E \sim 20 \text{ meV} \). This band also exhibits higher intensity at large momentum transfers, indicating lattice fluctuations as the origin. However, it should be noted that a peak in the aluminum phonon density of states exists near this energy transfer, thus suggesting that this particular band likely corresponds to scattering from the sample can and walls of the cryostat. At higher-energy transfers, two distinct bands of excitations are present over a large range of energy transfers spanning from \( \sim 30 \text{ to } 70 \text{ meV} \). The intensity of these bands weakens with momentum transfer \( L \); however, as illustrated in Fig. 4(e), these excitations overlap with lattice vibrational modes previously identified by infrared spectroscopy [106]. We will later discuss the origin of these excitations in terms of magnetovibrational scattering [115] by comparing the dispersion of these two high-energy bands at the magnetic zone boundary to those of phonons measured at large momentum transfers.

C. Comparison between experimental data and the spin-orbit exciton calculation

Having summarized our experimental data, we now present a direct comparison to our calculated parametrization based on the Green’s function approach that has been discussed above. Here, we first present the final model used to describe the experimental data. This is followed by a discussion concerning how such a model and its refined parameters were obtained.

As illustrated in Fig. 5, by allowing the value of mean molecular field \( H_{MF} \) to refine independently for each of the equally weighted 16 \( xAXxy \) domains, each possessing identical refined values of \( \lambda, J, \) and \( \Gamma \), our mean-field multilevel spin-orbit exciton model successfully reproduces both the fine structure at the magnetic zone center and the broad
FIG. 5. Comparison of \((\mathbf{Q}, E)\) slices and corresponding \(\mathbf{Q}\)-integrated cuts for CoO measured on MERLIN at 5 K and calculated with a mean-field multilevel spin-orbit exciton model employing the refined parameters listed in Tables III and II for an \(E_i\) of [(a), (d), (g)] 45 meV, [(b), (e), (h)] 110 meV, and [(c), (f), (i)] 75 meV. Horizontal bars in \(\mathbf{Q}\)-integrated cuts indicate experimental resolution. All \((\mathbf{Q}, E)\) slices have been folded along [001]. Individual contributions for each \(x\text{A}xy\gamma\) orbital configuration to the \(\mathbf{Q}\)-integrated cuts presented in (g) and (h) are illustrated in Fig. S1 [60].
excitations at the (1.5, 1.5, −1) zone boundary, while capturing the steeply dispersive columns of scattering observed at higher-energy transfers (Fig. 6). The refined values of the mean molecular field $H_{MF}$ are listed in Table II for the 16 $xAxyy$ orbital configurations, each with refined values of $\lambda$, $\Gamma$, and $J_{m,\xi}$ summarized in Table III.

The need for all 16 $xAxyy$ domains is illustrated in Figs. 7(e) and 7(f), where despite the success of the spin-orbit exciton model in reproducing the bandwidth of the excitations between $\sim 20$ meV up to $\sim 60$ meV with the initial parameters listed in Table I, the use of a single $xAxyy$ orbital configuration fails to reproduce the low-energy fine structure at the zone center and the steeply dispersive columns of high-energy scattering at the zone boundary, instead predicting the presence of a single dominant highly dispersive $G^{+}$ mode [Figs. 7(b) and 7(g)]. While both $G^{++}$ and $G^{--}$ modes exhibit negligible intensity [Fig. 7(d)], a direct consequence of the definition of $G^{\alpha\beta}$ given by Eq. (16) above, the spin-orbit exciton model does predict two additional gapped modes corresponding to the longitudinal $G^{zz}$ mode and the transverse $G^{++}$ mode. As summarized in Figs. 7(a) and 7(c), it is clear that both the weakly dispersive $G^{++}$ and $G^{zz}$ modes cannot account for the missing spectral weight in the fine structure.

**TABLE II.** Refined values (in meV) of the mean molecular field parameter $H_{MF}$ for all 16 $xAxyy$ orbital configurations considered in a mean-field multilevel spin-orbit exciton model, each with refined values of $\lambda$, $\Gamma$, and $J_{m,\xi}$ listed in Table III. Numbers in parentheses indicate statistical errors.

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<th>Orbital configuration</th>
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<tr>
<td>AAAA2</td>
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<tr>
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<td>Average</td>
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FIG. 7. A comparison of the calculated ($Q$, $E$) slices along (1.5, 1.5, $L$) for (a) $G^{+}$, (b) $G^{--}$, (c) $G^{zz}$, and (d) $G^{++}$ components of (e) the total response function $G$ and (f) the corresponding slice measured on MERLIN at $T = 5$ K with an $E_i = 110$ meV, with (g), (h) the same comparison for an $E_i = 45$ meV. The calculated model presented here only includes the AAAA orbital configuration. The model’s parameters’ values were fixed to those initial values listed in Table I. The ($Q$, $E$) slices presented in (f) and (h) have been folded along [001]. The intensity modulation observed in the calculated response for high-resolution measurements presented in (g) is an artifact of the steep dispersion of the excitation, manifesting itself as singularities in $g^{\alpha\beta}$ and creating numerical difficulties with sampling the $G^{+}$ mode.
as both modes are significantly weaker in intensity relative to the dominant $G^{\pm}$ mode, while contributing solely at higher-energy transfers ($\gtrsim 40$ meV). As illustrated in Figs. 6 and 8, while the longitudinal $G^O$ corresponds to a band of scattering that is broad in both momentum and energy while being located above the dominant $G^{+}$ component, the transverse $G^{-}$ corresponds to a flat band of scattering centered about $\sim 40$ meV.

In addition to the necessity for 16 $\Lambda \Lambda \Lambda \Lambda \gamma$ domains, the discrepancy between experiment and the calculated dispersion for each of the $G^{\pm}$ components that were presented in Fig. 7 confirmed the need for the optimization of the spin-orbit exciton model’s parameters. A summary of the influences of each of the model’s parameters for a fixed orbital configuration $J$ via $\lambda$, $H_{MF}$, and $\Gamma$ on the calculated spectra is presented in Fig. 9. Calculations are shown by false color maps and compared against constant momentum cuts at the $Q = (1.5, 1.5, −1.0)$ magnetic zone boundary and $Q = (1.5, 1.5, −0.5)$ zone center. For illustrative purposes, two different orbital configurations AAAA and AAAF are shown being denoted by solid and dashed lines, respectively.

As was previously noted in the discussion of the interion Hamiltonian $H_2$, the conversion factor $\hat{a}$ (Fig. 2) between the energy transfers measured by neutron spectroscopy and the magnitude of the corresponding magnetic exchange constants were determined by diagonalizing the pair Hamiltonian $H_{pair}$ [Eq. (11)]. Since deviations away from the linear dependence predicted by the projection theorem occur when the value of $|J| \rightarrow |\lambda|$, the value of larger magnetic exchange constants, such as the strong 180° antiferromagnetic superexchange $J_2$, is particularly sensitive to $|\lambda|$. As illustrated in Fig. 9(a), the strong sensitivity of $|J_2|$ on the value of $|\lambda|$ is of particular concern for CoO since the experimentally determined value of $\lambda = −16(3)$ meV possesses a significant relative error of almost 20% [54], corresponding to a large range of possible exchange values.

The influence of the large relative error for $\lambda$ is summarized in Fig. 9(b), illustrating that an increase of $|\lambda|$ by 20% to −19 meV from its initial value of −16 meV for a fixed value of a given molecular field $H_{MF}$ results in a significant shift to higher-energy transfers for the dominant $G^{+}$ component at both the magnetic zone center and boundary. This increase in energy transfer is much more prominent for the $G^{+}$ component compared to the less intense $G^{\pm}$, with the greatest increase for both components occurring at the magnetic zone center. Similar behavior in the calculated response is observed for a change in the mean molecular field $H_{MF}$ as illustrated in Fig. 9(c), with shifts in energy transfers significantly larger compared to the same relative change in the value of $|\lambda|$. In contrast with the three parameters presented so far, the influence of the tetragonal distortion parameter $\Gamma$ is most pronounced on the less intense weakly dispersive $G^{+}$ mode. As illustrated in Fig. 9(d), $\Gamma$ provides a mechanism to shift the energy transfers of the $G^{+}$ mode without inducing a comparable shift for the dominant $G^{-}$ mode for a fixed set of values for $H_{MF}$ and $J$ via a fixed value for $\lambda$.

With such a large number of domains under consideration, constraints on the parameter space for $J$ via $\lambda$, $\Gamma$, and $H_{MF}$ were required to ensure convergence for a least-squares optimization. As summarized in Table III, the model presented in Fig. 5 allowed $\lambda$ to vary from $−13$ to $−19$ meV, corresponding to the experimental error associated with its reported value [54], while the value of $\Gamma$ varied from $−8.76$ to $−6.16$ meV, corresponding to the range of its values reported in the literature [54]. The exchange constants $J_{m,\xi}$ for each coordination shell $m = 1 \ldots 4$, and type $\xi = A$ and $F$ were allowed to vary $\pm 20\%$ from their starting values obtained from the excitation energies derived from Mg0.97Co0.03O [55] with a Hamiltonian $H_{pair}$ employing the value of $\lambda$ under consideration. Such a deviation of the exchange constants was rationalized by noting that the estimates of the exchange constants listed in Table I are based on the superexchange pathways present in Mg0.97Co0.03O. Due to the prevalence of the nonmagnetic

<table>
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<td>$H_{MF}$</td>
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<td>$[0, 100]$</td>
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FIG. 8. A comparison of $(Q, E)$ slices (a) measured at 5 K on MERLIN and (b) calculated using spin-orbit exciton model for an $E_z = 110$ meV. (c) Comparison between (a) and (b) in a corresponding $Q$-integrated cut. The success of the spin-orbit exciton model to reproduce the $G^{−}$ component is emphasized. $(Q, E)$ slices presented in (a) and (b) have been folded along [001].

TABLE III. Summary of the initial values, parameter spaces, and refined values for the parameters of the mean-field multilevel spin-orbit exciton model. All values are reported in meV and numbers in parentheses indicate statistical errors.
FIG. 9. (a) Influence of the spin-orbit exchange constant on the splitting of the $J_{\alpha\alpha} = \frac{1}{2}$ manifold as a function of $|J|$ for the pair Hamiltonian $\hat{H}_{\text{pair}}$ [Eq. (11)]. Co$^{2+}$ pair excitations previously measured on MARI are shown explicitly with the corresponding value of $J_2$ (inset) particularly influenced by the nonlinearity at high-energy transfers. Calculated $(Q,E)$ slices for $E_i = 110$ and 45 meV with corresponding $Q$-integrated cuts at the zone boundary and center, respectively, illustrating the influences of (b) the spin-orbit coupling constant $\lambda$, (c) the mean molecular field $H_{\text{MF}}$, and (d) the tetragonal distortion $\Gamma_1$ parameters. Solid and dashed lines denote AAAA and AAAF orbital configurations, respectively. The prominent $G^{-+}$ and much weaker $G^{+-}$ have been labeled explicitly for reference. Unless otherwise stated, the values of $\lambda$, $H_{\text{MF}}$, $\Gamma_1$, and magnetic exchange constants $J$ were set to their initial values described in the main text and listed in Table I.

Mg$^{2+}$ in the dilute monoxide, these pathways and their superexchange constants most likely differ compared to those present in pure CoO. As a first approximation, the relative deviation of all magnetic exchange constants $J$ from their respective values in Mg$_{0.97}$Co$_{0.03}$O were set to be equal for all coordination shells $m$ and type $\xi$. As was previously discussed in Sec. IID, the values of $\lambda$, $J$, and $\Gamma$ were constrained to be equal for each of the 16 domains, while no such restriction was applied to the mean molecular field $H_{\text{MF}}$ which was itself allowed to vary independently for each domain from a value of 0 to an arbitrarily large upper limit. In our model, this limit was set to 100 meV, corresponding to $\sim$1.5 times the value of the initial value of 64.8 meV.

Among all the spin-orbit exciton model’s parameters, the expansion of the parameter space for the exchange constants $J_{m,\xi}$ proved to be of particular importance for the model’s
success in reproducing both the fine structure at the magnetic zone center and the broad excitations at the (1.5, 1.5, −1) zone boundary simultaneously. As summarized in Sec. VI of the Supplemental Material [60], the restriction of the exchange constants $J_{\text{ex}}$ to be equal to their respective values originally reported for $\text{Mg}_0.97\text{Co}_{0.03}\text{O}$ [55] for a given value of $\lambda$ resulted in failure of the spin-orbit exciton model to reproduce both broad excitations at the (1.5, 1.5, −1) zone boundary (Fig. S2 [60]). As summarized by Table III [60], such a restriction placed on the values of exchange constants resulted in the value of $\lambda$ being refined to its most negative permissible value. Such behavior is a reflection of the model’s attempts to minimize the average value for $H_{\text{MF}}$ in order to capture the intensity at lower-energy transfers at the (1.5, 1.5, $L$) zone boundary. As illustrated in Fig. S3 [60], the clear failure of the model, even by expanding the parameter space of $\lambda$, to include all values down to $-23$ meV, confirmed the inability of the model to reproduce the data at the zone boundary while employing the exact exchange constants measured in $\text{Mg}_0.97\text{Co}_{0.03}\text{O}$.

D. Magnetovibrational scattering and the magnetic zone boundary

Despite the success of the spin-orbit exciton parametrization of the experimental data along (1.5, 1.5, $L$), the model still fails at the (2, 2, $L$) zone boundary, suggesting the presence of additional physics that is beyond our localized model. The presence of such prominent magnetic scattering near (2, 2, $L$) is particularly unusual due to the predictions from both our spin-orbit exciton model and linear spin-wave theory for nearly zero intensity for magnetic fluctuations at these points in reciprocal space for an antiferromagnetic structure.

By comparing the scattering along (2, 2, $L$) to phonon branches near different nuclear zone boundaries, we note that there is a distinct overlap in dispersion. As is summarized in Fig. 10(a), a strong and steeply dispersing optical phonon mode near $Q = (5.5, 3.5, 4.5)$ exhibits an identical dispersion to that of the mode centered about (2,2,0), a mode that is not accounted for by our localized model. The phonon nature of the gapped mode in Fig. 10(a) is confirmed by the fact that the intensity increases with $Q$. A similar observation is illustrated in Fig. 10(c) for energy transfers greater than 60 meV, where a gapped phonon mode, identified by both its $Q$ dependence and previously reported first-principles calculations [113], exhibits a dispersion that is identical to the magnetic scattering around (2,2,0).

The apparent similarity between the dispersion of phonons at high $Q$ and the magnetic scattering that was originally unaccounted for in our model may suggest that our data at the zone boundaries include a magnetovibrational contribution to the neutron cross section. Corresponding to an indirect energy exchange between the neutron and a vibrating nucleus via the electromagnetic interaction between the neutron and the unpaired electrons of the nucleus, the magnetovibrational neutron cross section [115] is identical to the more commonly employed direct one-phonon cross section employing the nuclear force, with the exception that the nuclear scattering length $b$ is replaced by the magnetic scattering length given by

$$b_{\text{mag}} = \frac{e^2\gamma}{mc^2} f(Q) |\mu| \sin \alpha,$$

where $\alpha$ is the angle between the momentum transfer $Q$ and the ordered magnetic moment direction $\mu$. The presence of the isotropic magnetic form factor $f(Q)$ guarantees that the cross section will ultimately decay with momentum transfer, regardless of the underlying phonon origin of the scattering. This particular cross section requires an ordered magnetic structure and has been used previously to characterize the dynamic magnetic form factor [116]. Similar cross sections
have been reported in the rare-earth magnetic pyrochlores [117] and doped manganites [118]. We note that in the case of CoO, the magnetic ions are strongly correlated and coupled in all three dimensions, implying that the correlations are maintained at large energy transfers in the range of 30–60 meV where prominent phonon modes exist. Based on the comparison presented above, we suggest that both the excitations along (2, 2, L), and other excitations that were unaccounted for by our model such as the weak magnetic scattering at \( \sim 40 \) meV at the \((1.5, 1.5, L)\) zone boundary all originate from phonon modes, rather than the underlying magnetic Hamiltonian \( \hat{H} \).

While we have cast this discussion in terms of the magnetochemical cross section, which itself does not provide any new information on the underlying Hamiltonian, these excitations may be indicative of a coupling between the lattice and magnetic degrees of freedom. Such magnetoelastic coupling in the underlying Hamiltonian has been investigated recently in CeAuAl\(_3\) [119]. It is particularly compelling to consider such a claim given that these modes are located near the expected energy scale for the single-ion spin-orbit transitions from \( j_{\text{eff}} = \frac{5}{2} \rightarrow j_{\text{eff}} = \frac{3}{2} \). We will not investigate this coupling further given the need for first-principles calculations to reconcile past reported phonon data and the modes identified in our current study.

**E. High-energy response and the Co\(^{2+}\) form factor**

We have demonstrated that the experimental data at low-energy transfers can be successfully parametrized in terms of spin-orbit excitons with crystal field and exchange parameters based on our previous work reported on MgO substituted with Co\(^{2+}\). In Sec. III C, such a mean-field multilevel spin-orbit exciton model was shown to successfully reproduce the data in pure CoO near the zone center for energy transfers below 40 meV, while the failure of the model near the magnetic zone boundaries up to energy transfers of \( \sim 60 \) meV was attributed in Sec. III D to either magnetovibrational scattering or a coupling to underlying phonon excitations that were identified at large momentum transfers. In this final subsection, we will address the magnetic excitations at large energy transfers above \( \sim 65 \) meV, where no phonon scattering is expected, exceeding the dynamic range predicted by first-principles calculations [113] and measured by both our time-of-flight experiment and previous experiments [53,114,120].

In Fig. 11 we present a comparison between the scattering for high-energy transfers \((70, 105)\) meV with the lower-energy magnetic fluctuations that were discussed above in terms of our spin-orbit exciton parametrization. As summarized in Fig. 11(a), excitations located at low momentum transfers extend up to high energies. Previously identified in Fig. 6 as being magnetic in origin, it appeared that these excitations were successfully reproduced by the spin-orbit exciton model. These “columns” of scattering were interpreted as a result of the overlap at the magnetic zone boundary of multiple \( G^{-+} \) components from different local orbital arrangements that were required to capture the fine structure of the strongly dispersing fluctuations [Fig. 11(b)] centered about the zone centers and found at lower-energy transfers. While the analysis presented so far suggests that the \( Q \) dependence for all magnetic excitations appears to follow the Co\(^{2+}\) form factor, the energy-integrated slice in Fig. 11(a) demonstrates this is not the case for high-energy transfers, where the intensity decays more rapidly than \( f^2(Q) \). Such an observation can be confirmed by comparing momentum cuts of the magnetic fluctuations at small momentum transfers with the magnetic form factor for Co\(^{2+}\). In contrast with the lowest-energy transfers \([20–40]\) meV, Fig. 11(e) where the magnetic
fluctuations follow the form factor, deviations from such localized magnetic behavior begin to appear at (1.5, 1.5, ±0.5) for intermediate energy transfers [[40,70] meV, Fig. 11(d)], with the deviations being particularly prominent around (1.5,1.5,0) at the highest-energy transfers [[70,105] meV, Fig. 11(e)]. The rapid decay of intensity with momentum transfer may be indicative of a real-space object that is extended spatially, and not due to localized magnetism.

We now speculate as to the possible origin for such delocalized magnetism. In the case of the highest-energy transfers in CoO, where the relative deviation from the form factor is greatest, the scattering is steeply “dispersive” in energy, indicative of a large underlying energy scale. These fluctuations are highly reminiscent of the magnetic response found in itinerant magnets such as CeRhIn₅ [121], the cuprates [16,17], and also iron-based systems [122,123]. In the case of CeRhIn₅, the high-energy steeply dispersive excitations were found to be longitudinally polarized and occupied a region in (Q, E) phase space where two magnon processes were kinematically allowed. Termed the “1 + 2” model, multimagnon decay processes were used to provide a heuristic description of the data in the case of CeRhIn₅. Motivated by the qualitative similarities in the excitation spectrum between CeRhIn₅ and CoO, we investigated the possibility that the steeply dispersing excitations observed in CoO overlap a similarly allowed region of (Q, E) phase space. The phase space permitted for such a decay of the low-energy magnetic fluctuations was calculated using a simple model based on energy and momentum conservation that is given by [124–126]

\[
G(Q, E) = \sum_{Q_1, Q_2} \delta(Q - Q_1 - Q_2) \delta(E - E_{Q_1} - E_{Q_2}),
\]

where \(E_{Q_i}\) are the energies of transverse excitations at a given momentum transfer. As shown in Fig. 12, the kinematically allowed region overlaps in both momentum and energy with the steeply dispersing excitations at the highest-energy transfers; a region where the spin-orbit exciton model predicts the presence of a longitudinally polarized excitations (i.e., the \(G^{\text{CO}}\) mode), as was previously illustrated in Fig. 6(b).

Employing the observed overlap that is summarized in Fig. 12, we speculate that the spectral weight for these steeply dispersive high-energy excitations that are localized in momentum draws from this longitudinally polarized mode, analogous to what was observed in previously investigated itinerant magnets. Magnetism spatially extending beyond the Co²⁺ site has been suggested theoretically with some moment expected to be presented on the oxygen atom [32]. Given the extended nature of the magnetism in such a scenario, the magnetism would be expected to decay faster in momentum transfer than the isotropic Co²⁺ form factor. An intermediate example has been reported in the case of Sr₂CuO₄ [127] which found excellent agreement between the magnetic form factor and a model including strong covalent bonding and hybridization of the 3d orbitals. It could be that the higher-energy excitations are more sensitive to such a situation in CoO.

IV. CONCLUDING REMARKS

We have presented a neutron spectroscopic study of the magnetic fluctuations in the Mott insulator CoO. We have parametrized the low-energy magnetic excitations near the magnetic zone center in terms of a mean-field multilevel spin-orbit exciton model incorporating multiple structural domains owing to multiple local orbital arrangements, as well as a prominent tetragonal Jahn-Teller distortion. Dispersive excitations at the zone boundaries mimicking magnetic scattering at low Q that were originally unaccounted for by the spin-orbit exciton model were found to exhibit similar dispersions as phonons measured at larger momentum transfers, suggesting that the model’s failures at the zone boundaries may be attributed to magnetovibrational scattering or possibly coupling to lattice degrees of freedom. Finally, we report a discrepancy between the Q dependence of excitations at high-energy transfers and the behavior predicted by the Co²⁺ form factor. Despite the strong insulating nature of CoO, we speculate that such a discrepancy corresponds to a breakdown of spin-orbit excitons that may be accompanied by a crossover from localized to spatially extended magnetism, reminiscent of an itinerantlike response or strong covalency.

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