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Nonlinear NMR and magnon BEC in antiferromagnetic materials with coupled electron and nuclear spin precession

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We present a study of nonlinear NMR and Bose-Einstein condensation (BEC) of nuclear spin waves in antiferromagnetic MnCO₃ with coupled electron and nuclear spins. In particular, we show that the observed behavior of NMR signals strongly contradicts the conventional description of paramagnetic ensembles of noninteracting spins based on the phenomenological Bloch equations. We present a theoretical description of the coupled electron-nuclear spin precession, which takes into account an indirect relaxation of nuclear spins via the electron subsystem. We show that the magnitude of the nuclear magnetization is conserved for arbitrary large excitation powers, which is drastically different from the conventional heating scenario derived from the Bloch equations. This provides strong evidence that the coherent precession of macroscopic nuclear magnetization observed experimentally can be identified with the BEC of nuclear spin waves with $\mathbf{k} = 0$.

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I. INTRODUCTION

Bose-Einstein condensation (BEC) of bosons, or the accumulation of a macroscopic number of bosons in the same quantum state, presents one of the cornerstone subjects of studies in modern condensed-matter physics. In addition to the well-known phenomena of superfluidity, superconductivity, and BEC of cold atoms, the BEC of spin waves (magnons) in magnetically ordered materials has attracted a lot of recent attention [1-3]. Unlike conventional atomic BEC obtained by cooling atomic systems, magnon BEC can be established, for example, by continuous external rf pumping, which compensates for the loss of quasiparticles. The uniform precession of ordered spins in such systems can be described by magnons in a single quantum state with the wave number k = 0, which thus can be called the BEC of magnons in the $\mathbf{k} = 0$ state. On the other hand, a qualitatively similar state of uniform precession exists for a paramagnetic system of noninteracting spins under resonant excitation by an external rotating magnetic field. Since such a system of noninteracting spins cannot be described in terms of magnons, it is important to establish a clear physical picture of the BEC of magnons with k = 0 in magnetically ordered systems, and to distinguish it from the uniform precession of noninteracting spins induced by external rf excitation.

In this paper, we focus on magnetically ordered spin systems with coupled electron-nuclear spin precession. In such

systems, the dynamics of paramagnetic nuclear spins is governed by the exceptionally strong (10^5-10^6 Oe) hyperfine field from ordered electron spins. In turn, the motion of electron spins is affected by the hyperfine field from nuclear spins. This results in coupled electron-nuclear spin oscillations that can be described in terms of electron and nuclear spin waves [4,5]. In particular, a hyperfine interaction leads to the hybridization of, on the one hand, the usual spin waves in the magnetically ordered electron subsystem, i.e., *e*-magnons, and, on the other hand, the precession of a nuclear spin around the direction of the hyperfine field. Figure 1 shows examples of spectra for hybridized *e*-magnons (frequency Ω_+) and *n*-magnons (frequency Ω_-) in a weakly anisotropic antiferromagnet, MnCO₃. At zero wave number, k = 0, the frequency of *n*-magnons to a good approximation can be written as [6]

$$\Omega_{-}(0) \approx \omega_n - \omega_p(H, T) \cos \beta, \tag{1}$$

where ω_n is the frequency of precession of the nuclear spin in a stationary hyperfine field (that is, when the oscillations of the electron spin system caused by coupling to the nuclear spins are ignored), ω_p is a quantity that depends on a number of parameters including the applied magnetic field H and temperature of the nuclear system T, and β is the angle of deflection of the nuclear magnetization vector from its equilibrium orientation \mathbf{z} . The second term on the right-hand side of (1) is usually known as "frequency pulling." It is clear that the frequency $\Omega_-(0)$ corresponds to the usual NMR frequency, which can be measured in the experiment by applying a uniform ac magnetic field $\mathbf{h}(t) \perp \mathbf{z}$. Note that at large k the spectrum of the nuclear branch approaches ω_n . Therefore, for the existence of *n*-magnons, the frequency difference between $\Omega_-(0)$ and

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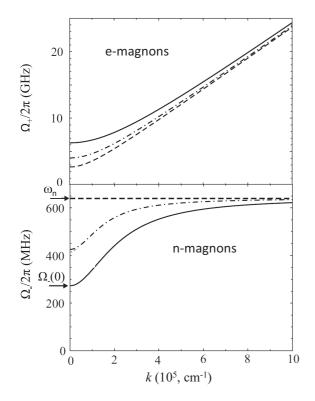


FIG. 1. The spectra of *e*-magnons and *n*-magnons in $MnCO_3$ in the static magnetic field of 0.2 kOe and at two values of temperature of 5 K (dashed-point line) and 1.4 K (solid line). The dashed lines show corresponding spectra in the case when coupling between electron and nuclear oscillations is neglected.

 ω_n must satisfy the condition

$$\omega_n - \Omega_-(0) \gg \delta \omega_n, \tag{2}$$

where $\delta \omega_n$ is the spread in the frequency ω_n due to inhomogeneities of the atomic and magnetic structures of the sample (inhomogeneous broadening) and due to relaxation processes (homogeneous broadening). There are few crystalline materials that satisfy this condition. Most notable examples are the weakly anisotropic antiferromagnets of the easy-plane type with ⁵⁵Mn as the magnetic ions, such as MnCO₃, CsMnF₃, RbMnF₃, and KMnF₃.

The existence of the *n*-magnon branch with minima at k = 0(see Fig. 1) suggests the possibility for the BEC of nuclear magnons with k = 0 under external rf pumping. Recent investigations in easy-plane, antiferromagnetic CsMnF₃ and MnCO₃ crystals using cw and pulsed NMR methods have shown evidence of magnon BEC [7-9]. In particular, the relatively strong and long-lived nuclear precession signals observed in pulsed NMR experiments indicated that nuclear magnetization is deflected by a large angle from the equilibrium orientation, while preserving its magnitude. This would strongly contradict the conventional behavior of a paramagnetic ensemble of noninteracting nuclear spins under external rf pumping, which is traditionally described by the phenomenological Bloch equations and predicts a strong reduction of the magnitude of the nuclear magnetization vector due to heating of the nuclear spin system. Thus, a detailed study and a more adequate description of coupled electron-nuclear spin systems under external rf pumping is necessary to establish a clear physical picture of magnon BEC in such systems.

Here, we present a detailed study of coupled electronnuclear spin dynamics in easy-plane, weakly anisotropic, antiferromagnet MnCO₃, under continuous NMR excitation. Our theoretical analysis shows that the coupling between electron and nuclear spin precession provides a mechanism not only for correlations between nuclear spins, that is, the formation of *n*-magnons, but also relaxation of nuclear spins via the electron subsystem. This mechanism of indirect relaxation preserves the magnitude of the nuclear magnetization vector for arbitrary large excitation powers; thus, it excludes the possibility of heating the nuclear subsystem by external pumping. This theory accounts well for the behavior of nonlinear NMR signals obtained in an antiferromagnetic MnCO3 sample at temperatures below 1 K. It is also shown that the conventional Bloch approach fails to account for our experimental results. Finally, we show that in the absence of heating, the energy of external pumping can be entirely transferred into the energy of uniform precession of nuclear spins, and the latter can be associated with the BEC of *n*-magnons with $\mathbf{k} = 0$.

In Sec. II, we provide a macroscopic description of coupled electron-nuclear spin systems based on equations of motion linearized with respect to transverse components of the electron magnetization vector. In Sec. III, we describe measurements of nonlinear NMR signals in an antiferromagnetic $MnCO_3$ sample and provide a detailed comparison with our theory. In Sec. IV, we discuss a microscopic picture of coupled electron-nuclear spin systems under excitation in terms of *n*-magnons and the formation of a uniform precession of nuclear spins, which can be identified as magnon BEC. Some details for the derivation of the coupled equations of motion for an electron-nuclear spin system are given in the Appendix.

II. THEORETICAL DESCRIPTION

Our theoretical analysis of a coupled electron-nuclear spin system in easy-plane, antiferromagnet $MnCO_3$ is based on the classical Landau-Lifshitz-Gilbert equations for macroscopic magnetization vectors of magnetic sublattices [10–14]. As usual, such an analysis of quantum spin systems is justified because the quantum-mechanical equations of motion reduce to classical equations for the precession of macroscopic magnetization vectors after averaging over the quantum-mechanical density matrix operator. Following the previous work [15], we describe the dynamics of a coupled electron-nuclear spin system using equations for two electron (M_1 and M_2) and two nuclear (m_1 and m_2) magnetization vectors of two sublattices,

$$\frac{d\mathbf{M}_{1}}{dt} = \gamma_{e} (\mathbf{M}_{1} \times \mathbf{H}_{M_{1}}) + \mathbf{R}_{M_{1}},$$

$$\frac{d\mathbf{M}_{2}}{dt} = \gamma_{e} (\mathbf{M}_{2} \times \mathbf{H}_{M_{2}}) + \mathbf{R}_{M_{2}},$$

$$\frac{d\mathbf{m}_{1}}{dt} = \gamma_{n} (\mathbf{m}_{1} \times \mathbf{H}_{m_{1}}) + \mathbf{R}_{m_{1}},$$

$$\frac{d\mathbf{m}_{2}}{dt} = \gamma_{n} (\mathbf{m}_{2} \times \mathbf{H}_{m_{2}}) + \mathbf{R}_{m_{2}},$$
(4)

where \mathbf{H}_{M_1} and \mathbf{H}_{M_2} (\mathbf{H}_{m_1} and \mathbf{H}_{m_2}) are the effective magnetic fields acting on each electron (nuclear) sublattice, and \mathbf{R}_{M_1}

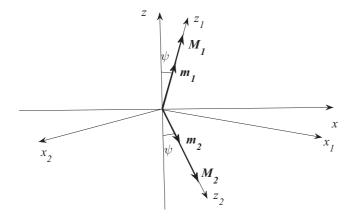


FIG. 2. Equilibrium position of the electron and nuclear magnetization vectors for each sublattice.

and \mathbf{R}_{M_2} (\mathbf{R}_{m_1} and \mathbf{R}_{m_2}) are the relaxation terms for the electron (nuclear) magnetization in each sublattice. The effective magnetic fields are given by functional derivatives

$$\mathbf{H}_{\mathbf{M}_{i}} = -\frac{\delta\Phi}{\delta\mathbf{M}_{i}}, \quad \mathbf{H}_{\mathbf{m}_{i}} = -\frac{\delta\Phi}{\delta\mathbf{m}_{i}}, \quad i = 1, 2.$$
(5)

The thermodynamic potential of the whole spin system, including its interaction with external static and rf magnetic fields, is given by

$$\Phi = \Phi_{\text{ex}} + \Phi_A + \Phi_D + \Phi_H + \Phi_{\text{hyp}}.$$
 (6)

The exchange interaction between electron spins of the two sublattices is described by

$$\Phi_{\rm ex} = \frac{1}{V} \int d\mathbf{r} \int d\mathbf{r}' J(r-r') \mathbf{M}_1(r) \mathbf{M}_2(r'), \qquad (7)$$

where J > 0 describes the exchange interaction and V is the sample volume. The energy of magnetic anisotropy with a dedicated axis **c** chosen as the positive **y** direction is described by

$$\Phi_A = \frac{1}{V} \int d\mathbf{r} K \{ \left[M_1^y(r) \right]^2 + \left[M_2^y(r) \right]^2 \}.$$
(8)

In the antiferromagnet MnCO₃ considered here, this anisotropy is the easy plane that corresponds to the condition K > 0. The Dzyaloshinskii-Moriya interaction which is responsible for the noncollinearity of the \mathbf{M}_1 and \mathbf{M}_2 sublattices (see Fig. 2) is described by

$$\Phi_D = \frac{1}{V} \int d\mathbf{r} D \Big[M_1^x(r) M_2^z(r) - M_1^z(r) M_2^x(r) \Big], \quad (9)$$

where D is a constant that characterizes the strength of the Dzyaloshinskii-Moriya interaction. The energy of \mathbf{M}_1 and \mathbf{M}_2 magnetizations in the external static magnetic field \mathbf{H} applied in the positive \mathbf{x} direction and the ac magnetic field $\mathbf{h}(t)$ applied along axis z is described by

$$\Phi_{H} = -\frac{1}{V} \int d\mathbf{r} H \Big[M_{1}^{x}(r) + M_{2}^{x}(r) \Big] -\frac{1}{V} \int d\mathbf{r} h(t) \Big[M_{1}^{z}(r) + M_{2}^{z}(r) \Big].$$
(10)

In what follows, we neglect the direct interaction of nuclear magnetization vectors \mathbf{m}_1 and \mathbf{m}_2 with the static and rf magnetic fields compared to the much stronger hyperfine field. However, note that the rf magnetic field still strongly couples to the nuclear spins via the $\mathbf{h}(t)$ -induced oscillations of electron magnetization. The hyperfine interaction between electron and nuclear spins, which for simplicity we assume to be isotropic, is described by

$$\Phi_{\text{hyp}} = -\frac{1}{V} \int d\mathbf{r} A[\mathbf{M}_1(r)\mathbf{m}_1(r) + \mathbf{M}_2(r)\mathbf{m}_2(r)]. \quad (11)$$

At $\mathbf{h}(t) = 0$, the minimum of the functional (6) defines the equilibrium orientation of the vectors $\mathbf{M}_{1,2}$ and $\mathbf{m}_{1,2}$ (see Fig. 2). The small angle ψ in this figure is determined from the equation

$$\sin\psi = \frac{H + H_D}{H_E},\tag{12}$$

where $H_D = DM_0$ is the magnitude of the Dzyaloshinskii-Moriya field, $M_0 = |\mathbf{M}_1| = |\mathbf{M}_2|$ is the magnitude of the electron magnetization vector in each sublattice, and the exchange field is given by

$$H_E = 2M_0 \int d(\mathbf{r} - \mathbf{r}') J(\mathbf{r} - \mathbf{r}').$$
(13)

The typical angle ψ is on the order of 10^{-2} rad.

The solutions of Eqs. (3)–(11) for small deviations of $\mathbf{M}_{1,2}$ and $\mathbf{m}_{1,2}$ from their equilibrium orientations were investigated earlier [4,15]. In this case, the eigenmodes of coupled electron-nuclear oscillations can be found from Eqs. (3) and (4) linearized with respect to the transverse components of magnetization vectors $\mathbf{M}_{1,2}$ and $\mathbf{m}_{1,2}$ and by taking the rf field $\mathbf{h}(t)$ and relaxation terms to be zero. As expected, the corresponding eigenfrequencies coincide with frequencies of coupled electron-nuclear spin waves at k = 0 [see Eq. (1)]. Electron and nuclear ac susceptibilities can be obtained from the same linearized equations by including the homogeneous rf magnetic field $\mathbf{h}(t)$ [6].

In what follows, we are interested in finding solutions of Eqs. (3)–(11) for arbitrarily large deviations of vectors $\mathbf{m}_{1,2}$ from their equilibrium orientations due to the rf magnetic field at frequencies close to the NMR resonance. Following the previous work [14], it is convenient to introduce a separate reference frame $x_1y_1z_1$ ($x_2y_2z_2$) for sublattice \mathbf{M}_1 (\mathbf{M}_2) obtained from the original frame xyz by a clockwise rotation around the z axis by angle ψ (by angle $\pi - \psi$) (see Fig. 2), and we chose variables according to

$$2M^{\xi} = M_1^{x_1} + M_2^{x_2}, \quad 2m^{\xi} = m_1^{x_1} + m_2^{x_2},$$

$$2M^{\eta} = M_1^{y_1} + M_2^{y_2}, \quad 2m^{\eta} = m_1^{y_1} + m_2^{y_2}, \quad (14)$$

$$2M^{\zeta} = M_1^{z_1} + M_2^{z_2}, \quad 2m^{\zeta} = m_1^{z_1} + m_2^{z_2},$$

where $M_i^{x_i}, M_i^{y_i}, M_i^{z_i}$ $(m_i^{x_i}, m_i^{y_i}, m_i^{z_i})$, i = 1, 2, are components of electron (nuclear) magnetization vectors in the corresponding frame. The solutions of Eqs. (3) and (4) in these variables can by found by linearizing Eqs. (3) with respect to the transverse components of the electron magnetization vectors, whose deviation from the equilibrium orientation is assumed to be small. This assumption is valid for sufficiently large magnetic fields $H > H_c$ [Eq. (A2)], where the oscillation of electron spins is only weakly affected by the hyperfine interaction with nuclei (see the discussion in the previous section). The details of its derivation are given in the Appendix. Finally, is is convenient to write the result in the coordinate system XYZ, where the components of the net nuclear magnetization m_X , m_Y , and m_Z are related to m^{ξ} , m^{η} , and m^{ζ} by

$$m^{\xi} = m_X \cos \omega t + m_Y \sin \omega t,$$

$$m^{\eta} = -m_X \sin \omega t + m_Y \cos \omega t,$$
 (15)

$$M^{\zeta} = m_Z,$$

which is somewhat similar to the usual rotating wave transformation used to describe traditional NMR [16].

By omitting the relaxation terms in Eqs. (3) and (4), equations for the net nuclear magnetization become (see the Appendix for details)

$$\frac{dm_X}{dt} = \left(\omega_n - \omega - \omega_p \frac{m_Z}{m_0}\right) m_Y,$$
(16)

$$\frac{dm_Y}{dt} = -\left(\omega_n - \omega - \omega_p \frac{m_Z}{m_0}\right) m_X + \omega_1 m_Z, \quad (17)$$

$$\frac{dm_Z}{dt} = -\omega_1 m_Y,\tag{18}$$

where ω_1 is proportional to the amplitude of the rf magnetic field \mathbf{h}_1 and ω_p is the parameter discussed previously [see Eqs. (A10) and (A12) in the Appendix for the precise definition of ω_1 and ω_p]. Since the relaxation terms were neglected, the above equations can be used to describe the dynamics of the magnetization vector \mathbf{m} only on time intervals t, which are small compared to the nuclear magnetic relaxation time. Note that as was discussed previously from the microscopic point of view, the presence of the frequency pulling term $\omega_p(m_Z/m_0)$ does not allow us to obtain large deflection angles of **m** from the equilibrium orientation using a short rf pulse, as can typically be done in conventional NMR (the Rabi flopping) [16]. Indeed, for typical experimental conditions we have $\omega_p \gg \omega_1$, therefore the orientation of the axis of rotation of vector **m** in the frame XYZ changes quickly to be nearly parallel to Z, after applying the rf pulse. However, as we show below, it is possible to obtain large angles of deflection by cw NMR. The description of the stationary state of the spin system under cw rf pumping requires accounting for the proper form of relaxation terms in Eqs. (3) and (4).

As discussed previously, the interaction between transverse components of the electron and nuclear magnetization vectors significantly changes the frequency of nuclear spin oscillations. Another consequence of this interaction is the appearance of the relaxation of the transverse component of vector **m**. Indirect relaxation of this type has been studied previously, but for relatively small amplitudes of **m**-vector oscillations when nonlinear phenomena were insignificant [15]. Here, we consider nonlinear equations of motion for nuclear magnetization taking into account its indirect relaxation via the electron spin system.

Typically, relaxation of the transverse component of magnetization occurs due to fluctuating magnetic fields acting on different spins, and to inhomogeneity of the external static magnetic field acting in the electron system. In what follows, we use an assumption that, because nuclear spins experience an enormously large hyperfine field, $H_e \sim 10^{5}-10^{6}$ Oe, we can neglect any other fields, including fluctuating magnetic fields responsible for the relaxation of vectors $\mathbf{M}_{1,2}$. We note that the approximation $\mathbf{R}_{m_{1,2}} = 0$ in Eqs. (4) conserves the magnitude of vectors $\mathbf{m}_{1,2}$ and the net magnetization vector \mathbf{m} . Thus, this approximation excludes the possibility of heating of the nuclear spin system by any processes, including its excitation by rf pumping.

Using this approximation together with the assumption of small deviations of electron magnetization vectors from their equilibrium orientations at $H > H_c$, equations of motion for the vector **m** become (see the Appendix for details)

$$\frac{dm_X}{dt} = \left(\omega_n - \omega - \omega_p \frac{m_Z}{m_0}\right) m_Y - \frac{m_Z}{m_0} \frac{m_X}{T_{2n}}, \quad (19)$$

$$\frac{dm_Y}{dt} = -\left(\omega_n - \omega - \omega_p \frac{m_Z}{m_0}\right) m_X + \omega_1 m_Z - \frac{m_Z}{m_0} \frac{m_Y}{T_{2n}},$$
(20)

$$\frac{dm_Z}{dt} = -\omega_1 m_Y - \frac{m_Z + m_0}{m_0} \frac{m_Z - m_0}{T_{2n}},$$
 (21)

where the nuclear transverse relaxation time T_{2n} is related to the electron transverse relaxation time T_2e by Eq. (A13) in the Appendix.

The cw NMR signals are described by stationary state solutions of the above equations. The latter are given by

$$\frac{m_X}{m_0} = \frac{(\omega_n - \omega - \omega_p + \omega_p \Delta)\omega_1(1 - \Delta)}{(\omega_n - \omega - \omega_p + \omega_p \Delta)^2 + \Gamma_1^2},$$
 (22)

$$\frac{m_Y}{m_0} = \frac{\omega_1 \Gamma_1 (1 - \Delta)}{(\omega_n - \omega - \omega_p + \omega_p \Delta)^2 + \Gamma_1^2},$$
 (23)

where $\Delta = 1 - m_Z/m_0$, which determines the longitudinal component of **m**, is the root of the fourth-degree polynomial equation

$$\Delta(2-\Delta) = \frac{\omega_1^2 (1-\Delta)^2}{(\omega_n - \omega - \omega_p + \omega_p \Delta)^2 + \Gamma_1^2},$$
 (24)

and Γ_1 is given by

$$\Gamma_1 = (1 - \Delta)/T_{2n}.$$
(25)

For the sake of qualitative comparison with our theory, let us also consider the traditional Bloch approach in which the nuclear relaxation is described by introducing two phenomenological relaxation times, i.e., the longitudinal magnetization relaxation time T_1 and the transverse magnetization relaxation time T_2 [16]. Adding the corresponding relaxation terms into Eqs. (16)–(18), we obtain the phenomenological Bloch equations

$$\frac{dm_X}{dt} = \left(\omega_n - \omega - \omega_p \frac{m_Z}{m_0}\right) m_Y - \frac{m_X}{T_2},$$
 (26)

$$\frac{dm_Y}{dt} = -\left(\omega_n - \omega - \omega_p \frac{m_Z}{m_0}\right)m_X + \omega_1 m_Z - \frac{m_Y}{T_2},\quad(27)$$

$$\frac{dm_Z}{dt} = -\omega_1 m_Y - \frac{(m_Z - m_0)}{T_1}.$$
 (28)

Note that the above equations are obtained by disregarding the electron spin relaxation terms $\mathbf{R}_{M_{1,2}}$ in Eqs. (3). The stationary state solutions of (26)–(28) can be written as

$$\frac{m_X}{m_0} = \frac{(\omega_n - \omega - \omega_p + \omega_p \Delta)\omega_1(1 - \Delta)}{(\omega_n - \omega - \omega_p + \omega_p \Delta)^2 + T_0^{-2}},$$
(29)

$$\frac{m_Y}{m_0} = \frac{\omega_1 T_2^{-1} (1 - \Delta)}{(\omega_n - \omega - \omega_p + \omega_p \Delta)^2 + T_2^{-2}},$$
(30)

where $\Delta = 1 - m_Z/m_0$, as defined earlier, is given by the roots of the third-degree polynomial equation

$$\Delta = \frac{\omega_1^2 T_1 T_2^{-1} (1 - \Delta)}{(\omega_n - \omega - \omega_p + \omega_p \Delta)^2 + T_2^{-2}}.$$
 (31)

To distinguish between the two approximations which lead, on the one hand, to Eqs. (22)–(24), and, on the other hand, to Eqs. (29)–(31), we need to compare the continuous NMR signals predicted by these equations with experimentally observed signals. This is done in the next section.

III. EXPERIMENTAL COMPARISONS

A. Experimental methods

In the experiment, we studied cw NMR signals in an easyplane antiferromagnetic MnCO₃ sample at temperatures below 1 K. The sample was in the form of a rhombus-shaped plate about 0.7 mm thick, with diagonals of approximately 2.4 and 2.7 mm. The magnetic anisotropy axis **c** was perpendicular to the plane of the plate. The measured mass of the sample was about 8×10^{-3} g.

To excite continuous NMR signals in MnCO₃ by external rf pumping, the sample was placed in a high-quality factor rf resonator of a split-ring type [17]. The resonator and sample were oriented such that the external static magnetic field H and the rf magnetic field h of the resonator were perpendicular to each other and to the anisotropy axis **c** of the sample. The resonator was made of high-conductivity, oxygen-free copper and had a resonant frequency of about 591.3 MHz and a quality factor of about 600 measured at 1 K. The input rf signal transmitted through a cryogenic, low-loss, semirigid coaxial cable was coupled to the resonator via a loop made at the end of the cable, and the reflected signal was measured at different values of static field H and input rf power P using a network analyzer. Signals proportional to the in-phase and quadrature (with respect to the rf field in the cavity) components of ac magnetization induced in the sample could be extracted from the reflected signal. In order to avoid complications associated with the strong coupling of the spin system to the electromagnetic mode of the resonator, which lead to normal mode splitting in the resonator reflection spectrum [18], we used the frequency of rf pumping of 593.5 MHz, which is significantly detuned from the resonant frequency of the resonator. In this so-called dispersive regime of coupling, the in-phase (M_d) and quadrature (M_a) components of the ac magnetization are related to the complex amplitude (phasor) of the reflected signal, according to

$$V \approx V_0 + C(-M_a + iM_d), \tag{32}$$

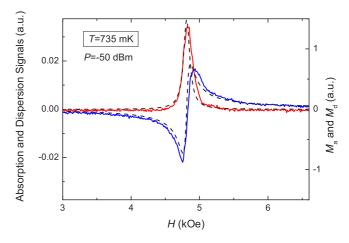


FIG. 3. Absorption (solid line, red) and dispersion (solid line, blue) signals measured in MnCO₃ at T = 735 mK and at a low input rf power of P = -50 dBm. Dashed lines are the ac magnetization components M_a and M_d calculated as described in Sec. III C.

where V_0 is the complex amplitude of the signal reflected from an empty cavity, *i* is the imaginary unit, and C > 0 is a proportionality coefficient that depends on the characteristics of the resonator and its coupling to the rf transmission line.

B. Experimental results

Figure 3 shows the NMR signals proportional to M_a (absorption signal) and M_d (dispersion signal) measured at T = 735 mK as a function of the static field H at a low input rf power of -50 dBm. At such low power, the deviation of nuclear magnetization from its equilibrium orientation is very small. Correspondingly, frequency pulling is very close to its value for the equilibrium spin system; therefore, nonlinearity of the NMR signal is negligible. These signals are used to find the fitting parameters that define the dependence of frequency pulling and the nuclear magnetic relaxation rate on the value of H. It should be noted that although these signals were measured in the linear regime of NMR, they are not ideally symmetric. This suggests that the sample used in the experiment is not a single crystal, which is also confirmed by careful visual examination under the microscope. It is likely that the sample consists of at least two or three monocrystals of different size with somewhat different relative orientations of their lattices.

Figures 4 and 5 show absorption and dispersion signals measured at T = 735 mK and at different input rf power levels. It is clear that with increasing power, the NMR frequency shifts toward lower values of H due to frequency pulling. Both absorption and dispersion signals become very asymmetrical. In addition, both signals show strong hysteresis upon reversing the direction of the field sweep. In Fig. 6 we show the absorption and dispersion signals obtained at the highest input power of P = 0 dBm used in this experiment. In the next section, we compare the measured absorption and dispersion signals with predictions of our theory [see Eqs. (22)–(24)]. In addition, we compare the experimental results with predictions based on the phenomenological Bloch equations [see Eqs. (29)–(31)].

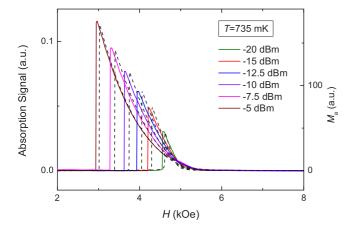


FIG. 4. Absorption signals (solid lines) measured in MnCO₃ at T = 735 mK and at several values of input rf power (shown in the figure) during a down-field sweep. Dashed lines represent values of M_a calculated as described in Sec. IIIC.

C. Expressions for ac magnetization components and their comparison with experimental results

In order to compare the results presented in Figs. 4 and 5 with predictions of our theory, first we need to find the relation between the ac magnetization components M_a and M_d and components of nuclear magnetization m_X and m_Y described by Eqs. (22) and (23). It is important to realize that due to the much larger magnitude of the electron magnetization vector $\mathbf{M} = (M^{\xi}, M^{\eta}, M^{\zeta})$ compared to that of the nuclear magnetization vector $\mathbf{M} = (m^{\xi}, m^{\eta}, m^{\zeta})$, the ratio of which is approximately $\gamma_e/\gamma_n \gtrsim 10^3$, the main contribution to the observed ac magnetization comes from oscillations of vector \mathbf{M} . We can find necessary expressions using the relation between the complex amplitudes of $M^{\xi}(t)$ and $m^{\xi}(t)$, as given by Eqs. (A8)–(A11) in the Appendix.

According to Fig. 2 and Eqs. (12)–(14), the vector component of electron magnetization $\mathbf{M}_{\parallel}(t) \parallel \mathbf{h}(t)$, which defines the

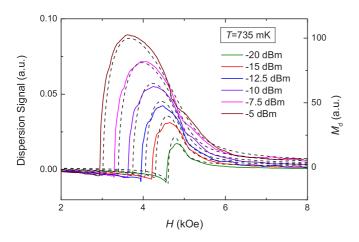


FIG. 5. Dispersion signals measured in MnCO₃ at T = 735 mK and at several values of input rf power (shown in the figure) during a down-field sweep. Dashed lines represent values of M_d calculated as described in Sec. III C.

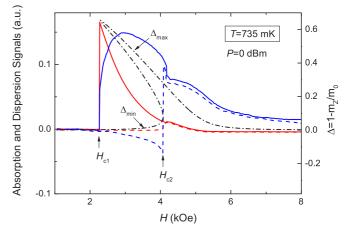


FIG. 6. Absorption and dispersion signals measured at T = 735 mK at the highest input rf power of P = 0 dBm during up-field (dashed lines) and down-field (solid lines) sweeps. The dashed-dotted line is the result of numerical calculations for $\Delta = 1 - m_Z/m_0$ as described in Sec. III C.

observed NMR signals, can be expressed in terms of $M^{\xi}(t)$ as

$$M(t) = M_1^{z} + M_2^{z} = -(M_1^{x_1} + M_2^{x_2}) \sin \psi$$

= $-2\left(\frac{H + H_D}{H_E}\right) M^{\xi}(t).$ (33)

Assuming time dependence of the rf magnetic field in the form $h(t) = 2H_1 \cos \omega t$, absorption M_a and dispersion M_d signals are given by, respectively, the imaginary and real parts of the complex amplitude of $M_{\parallel}(t)$. The expressions for M_d and M_a in terms of oscillating components of the nuclear magnetization vector can be found from the above equation using Eqs. (A8)–(A11) and approximations discussed in the Appendix. Using the transformation (15) and expressing results in terms of nuclear magnetization vector components m_X and m_Y , we obtain

$$M_{d} = M_{d}^{\mathrm{nr}} + \frac{H_{n}}{H} \left(1 - \frac{\omega^{2}}{\omega_{e}^{2}}\right)^{-1} \left[m_{X} - \frac{\omega}{\omega_{n}\omega_{p}T_{2n}}m_{Y}\right],$$
$$M_{a} = \frac{H_{n}}{H} \left(1 - \frac{\omega^{2}}{\omega_{e}^{2}}\right)^{-1} \left[m_{Y} + \frac{\omega}{\omega_{n}\omega_{p}T_{2n}}m_{X}\right].$$
(34)

In the above equations, the nonresonant contribution M_d^{nr} to the dispersion signal is given by

$$M_d^{\rm nr} = \frac{H_1 H_n}{H^2} \left(1 - \frac{\omega^2}{\omega_e^2} \right)^{-2} \frac{\omega_n}{\omega_p} m_0. \tag{35}$$

It arises from the direct coupling of the electron magnetization vector to the rf magnetic field having amplitude H_1 . The resonant contribution (at the NMR frequency) comes from the terms containing m_X and m_Y . The main contribution to the dispersion (absorption) signal comes from the $m_X (m_Y)$ component of the nuclear magnetization vector. Note that it is enhanced by the amplification factor η given by Eq. (A13) in the Appendix. The components m_X and m_Y are given by Eqs. (22) and (23) and are expressed in terms of the dimensionless parameter $\Delta = 1 - m_Z/m_0$ determined from Eq. (24). It is instructive to discuss the general form of the solutions of these equations. Equation (24) is a fourth-order polynomial equation with respect to Δ . One of the roots of Eq. (24), viz., $\Delta \approx 2$, corresponds to an unstable state with antiparallel orientation of the vectors **m** and **H**_n. The other three roots are the solutions of a third-order polynomial equation that is obtained from (24) by excluding the root $\Delta \approx 2$. We are interested in the real-valued roots and their dependence on *H*, which is defined by the corresponding dependencies of $\omega_1(H), \omega_p(H)$, and $T_{2n}(H)$ given by Eqs. (A12)–(A15) in the Appendix.

At low rf excitation powers, such that $\omega_1 T_{2n} \ll 1$, Eq. (24) has only one real root, given by

$$\Delta_1 = \frac{\omega_1^2}{(\omega_n - \omega - \omega_p)^2 + T_{2n}^{-2}} \ll 1.$$
(36)

Using this expression in Eqs. (22) and (23), we obtain

$$\frac{m_X}{m_0} = \frac{(\omega_n - \omega - \omega_p)\omega_1}{(\omega_n - \omega - \omega_p)^2 + T_{2n}^{-2}},$$
$$\frac{m_Y}{m_0} = \frac{\omega_1 T_{2n}^{-1}}{(\omega_n - \omega - \omega_p)^2 + T_{2n}^{-2}}.$$
(37)

At fixed rf frequency ω the position of the resonance in the static magnetic field *H* is determined by the dependence of ω_p on *H* given by Eq. (A14) in the Appendix. The resonance value of field $H = H_{\text{res}}$ is defined by

$$\omega_n - \omega - \omega_p(H_{\text{res}}) = 0. \tag{38}$$

The width of the resonance ΔH , which can be defined as the distance between the maximum and minimum of the dispersion signal, is related to T_{2n} as

$$T_{2n}^{-1}(H_{\rm res}) = -\frac{1}{2} \frac{\partial \omega_p(H_{\rm res})}{\partial H} \Delta H.$$
 (39)

The dependence of T_{2n} on H is given by Eq. (A15) in the Appendix. It is convenient to express the dependence of ω_p and T_{2n} on H in terms of adjustable parameters a_1 and a_2 as

$$\omega_p(H) = \frac{a_1}{H(H+H_D)},\tag{40}$$

$$T_{2n}^{-1}(H) = \frac{a_2}{H^2(H+H_D)^2},$$
(41)

and determine the numerical values of a_1 and a_2 at a given temperature from the comparison of experimental data taken at low rf excitation power and the ac magnetization components M_a and M_d calculated using Eqs. (34) and (37). Such a comparison is shown in Fig. 3, where M_a and M_d calculated using $a_1/2\pi = 2.06 \times 10^3$ MHz (kOe)² and $a_2/2\pi = 1.6 \times 10^3$ MHz (kOe)⁴ are plotted using dashed lines. Here, we used $H_D = 4.4$ kOe, $\omega_n/2\pi = 640$ MHz, and $\omega/2\pi = 593.5$ MHz. The value of H_1 was chosen to be sufficiently small to ensure a linear NMR regime. The corresponding values of $H_{\rm res}$ and ΔH are approximately 4.81 and 0.11 kOe, respectively. The width of the resonance corresponds to the nuclear relaxation T_{2n} time of approximately 0.19 μ s.

For arbitrarily large rf excitation powers there exists a range of H where there is more than one real-valued root of Eq. (24). The numerical solution of this equation for a large value of H_1 is shown by the dashed-dotted line in Fig. 6. In this calculation, we used $\omega_p(H)$ and $T_{2n}(H)$ given by Eqs. (40) and (41) with numerical values of a_1 and a_2 given above. The interval $H_{c1} < H < H_{c2}$ corresponds to the region where three real roots exist. The merger of two real roots (where they become complex) corresponds to the phenomenon which in mathematics is called *catastrophe*. We restrict ourselves to the analysis of the upper branch $\Delta_{\max}(H)$, which has its critical point at $H = H_{c1}$ (see Fig. 6). The state of the lower branch $\Delta_{\min}(H)$ is destroyed at fields $H < H_{c2}$ due to instabilities caused by the excitation of *n*-magnons that are in resonance with the rf pumping field [19,20].

In order to make a comparison between the experimental data presented in Figs. 4 and 5 and the ac magnetization components M_a and M_d calculated using Eqs. (22)–(24) and (34), we need to establish a correspondence between the values of the rf excitation power used in the experiment and the numerical values of H_1 used in the calculations. We used the value of $H_{c_1} \approx 2.2$ kOe in Fig. 6 to define the value of $H_1 =$ 23 Oe at the excitation power of 0 dBm. The line shapes of M_a and M_d calculated for the corresponding values of H_1 and using Eqs. (40) and (41) are plotted in Figs. 4 and 5, respectively, using dashed lines. As seen in these figures, we obtain very good quantitative agreement between our experimental data and predictions of our theoretical model based on Eqs. (19)-(21). Note that conservation of the magnitude of vector **m**, which follows from our theory, allows us to express $\Delta_{\max}(H)$ in terms of the deflection angle β_0 between the vector **m** and its equilibrium orientation,

$$\Delta_{\max}(H) = 1 - \cos \beta_0. \tag{42}$$

At the critical field $H_{c1} \approx 2.2$ kOe in Fig. 6 corresponding to the input rf power of 0 dBm, the deflection angle has a value $\beta_0 \approx 70^\circ$.

For the sake of comparison with the predictions of the Bloch model based on Eqs. (26)–(28), it is instructive to consider the dependence of the absorption signal M_a at $H = H_{c1}$ on the input rf power *P*. Note that the critical field $H = H_{c1}$ corresponds to the resonant condition

$$\omega_n - \omega - \omega_p + \omega_p \Delta = 0. \tag{43}$$

From Eqs. (22) and (23) for the steady state solutions, this gives $m_X(H_{c_1}) = 0$ and $m_Y(H_{c_1}) = m_0\omega_1T_{2n}$; therefore, from the second line of (34) and Eqs. (A12)–(A15) in the Appendix we obtain

$$M_a(H_{c_1}) = \frac{\gamma_n m_0 H_n^2}{a_2} \left(1 - \frac{\omega^2}{\omega_e^2}\right)^{-2} \left(H_{c_1} + H_D\right)^2 H_1. \quad (44)$$

In order to get rid of the field-independent factor in front of the right-hand side of the above equation, it is convenient to normalize this expression to the value of $M_a(H_{c1})$ at one of the values of the input rf power used in the experiment. For example, we choose the lowest power of $P^* = -20$ dBm used in Fig. 4. The expression for the normalized absorption signal becomes (for the sake of simplicity we neglect $\omega/\omega_e \ll 1$)

$$\frac{M_a(H_{c1})}{M_a^*(H_{c1})} \approx \left(\frac{H_{c1} + H_D}{H_{c1}^* + H_D}\right)^2 \frac{H_1}{H_1^*} = \alpha \left(H_{c_1}, \sqrt{P/P^*}\right), \quad (45)$$

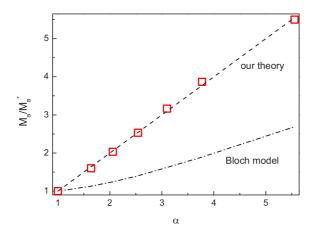


FIG. 7. Normalized values of the absorption signal at the critical field $H = H_{c1}$ as a function of quantity $\alpha(H_{c1}, \sqrt{P/P^*})$ defined by Eq. (45) obtained from the experimental data presented in Figs. 4 and 6 (open squares). The lines are the corresponding dependences predicted by our theory (dashed line) and by the conventional Bloch approach (dashed-dotted line).

where M_a^* , H_{c1}^* , and H_1^* are the corresponding values of M_a , H_{c1} , and H_1 at the input rf power $P = P^*$. Note that the normalized absorption signal depends only on the values of the critical field H_{c1} and the ratio $\sqrt{P/P^*} = H_1/H_1^*$, which can be readily determined in the experiment. Figure 7 shows the dependence of $M_a(H_{c1})/M_a^*(H_{c1})$ on $\alpha(H_{c1},\sqrt{P/P^*})$ obtained from the experimental data shown in Figs. 4 and 6 (opened squares), as well as the theoretically predicted dependence given by Eq. (45) (dashed line).

Now we compare this result with predictions of the heating scenario based on the phenomenological Bloch equations (26)–(28). Note that as in Eq. (24), the third-order polynomial equation (31) with respect to Δ can have three real-valued roots in certain ranges of the static magnetic field $H_{c1} < H < H_{c2}$. This may result in nonlinear NMR signals similar to those described earlier [4]. Using condition (43) satisfied at the field $H = H_{c1}$, we obtain from Eqs. (29)–(31),

$$m_x(H_{c1}) = 0, \quad m_y(H_{c1}) = \omega_1 T_2 m_z(H_{c1}),$$

$$m_Z(H_{c1}) = \frac{m_0}{1 + \omega_1^2 T_1 T_2}.$$
 (46)

Note that, unlike in our proposed theory, the Bloch model does not conserve the magnitude of the magnetization vector. Indeed, from Eq. (46), we have

$$|\mathbf{m}| = \sqrt{\frac{1 + \omega_1^2 T_2^2}{\left(1 + \omega_1^2 T_1 T_2\right)^2}} m_0.$$
(47)

Typically, we have $T_2 \ll T_1$; therefore, $|\mathbf{m}|$ can already be significantly less than m_0 for $\omega_1^2 T_1 T_2 \gtrsim 1$.

Using Eqs. (43) and (46), it is convenient to write $\omega_1^2 T_1 T_2$ as

$$\omega_1^2 T_1 T_2 = \frac{\Delta \omega(H_{c1})}{\omega_p(H_{c1}) - \Delta \omega(H_{c1})},\tag{48}$$

where $\Delta \omega = \omega - \omega_n + \omega_p$. Note that using the above equation, the values of $\omega_1^2 T_1 T_2$ at $H = H_{c1}$ can be determined

from the experimental values of H_{c_1} . Using the second line of Eq. (34) we obtain the expression for the absorption signal at the catastrophe field,

$$M_a(H_{c_1}) = \frac{m_0 H_n}{H} \sqrt{\frac{T_2}{T_1}} \left(1 - \frac{\omega^2}{\omega_e^2}\right)^{-1} \frac{\sqrt{\omega_1^2 T_1 T_2}}{1 + \omega_1^2 T_1 T_2}, \quad (49)$$

and using Eqs. (A12) and (A13) from the Appendix and the definition of α [see Eq. (45)], we obtain for the normalized absorption signal,

$$\frac{M_a(H_{c1})}{M_a^*(H_{c1})} \approx \left(\frac{H_{c1}^*(H_{c1}^* + H_D)}{H_{c1}(H_{c1} + H_D)}\right)^2 \frac{1 + \left(\omega_1^2 T_1 T_2\right)^*}{1 + \omega_1^2 T_1 T_2} \alpha, \quad (50)$$

where $(\omega_1^2 T_1 T_2)^*$ is the value of $\omega_1^2 T_1 T_2$ at the critical field for the input rf power $P = P^*$. Again, note that the term in front of α on the right-hand side of the above equation depends only on the value of the critical field $H = H_{c1}$, which can be determined from the experiment, and does not depend on the choice of values for T_1 and T_2 . The dependence of the normalized absorption signal on α given by Eq. (50) is plotted in Fig. 7 with a dashed-dotted line. Clearly, the conventional Bloch approach fails to adequately account for the behavior observed in the experiment. Using Eq. (48), it is straightforward to estimate the value of $\omega_1^2 T_1 T_2$ at the critical field $H = H_{c1}$ for the maximum input rf power P = 0 dBm. Using the value of $H_{c1} \approx 2.2$ kOe (see Fig. 6), we obtain $\omega_1^2 T_1 T_2 \approx 2$. Thus, the Bloch theory predicts a significant reduction of the magnitude of the nuclear magnetization vector due to heating. This does not agree with our experimental results.

IV. DISCUSSION

The mechanism of the indirect relaxation of nuclear spins via the electron subsystem, which preserves the magnitude of nuclear magnetization, has important consequences for the dynamics of *n*-magnons under external rf pumping. In this section we provide some detailed discussion of the processes involved and show that the pumping of *n*-magnons with a uniform ac magnetic field $\mathbf{h}(t)$ in a wide range of frequencies results in a macroscopic accumulation of *n*-magnons with $\mathbf{k} = 0$, which can be identified with the BEC of nonequilibrium *n*-magnons.

As pointed out earlier [20], an external rf pumping of the coupled electron-nuclear spin system with a uniform \mathbf{h} field at the frequency ω in the range $\Omega_{-}(0) < \omega < \omega_n$ can lead to instability against the formation of *n*-magnons with $k \neq 0$ corresponding to $\Omega_{-}(k) = \omega$ (see Fig. 8). The appearance of such magnons slightly decreases the deflection angle β of the nuclear magnetization vector from its equilibrium orientation. Further time evolution of such magnons depends on the nuclear relaxation processes. According to the Bloch approach scenario, such magnons thermalize and lead to a temperature increase of the nuclear system T. This would significantly decrease the magnitude of nuclear magnetization $m_0(T)$ and modify the spectrum of *n*-magnons such that the frequency of the k = 0 mode, given by Eq. (1), coincides with ω . A stationary state of the nuclear spin system would correspond to a small deflection angle $\beta = \arccos[m_z/m_0(T)]$. As we

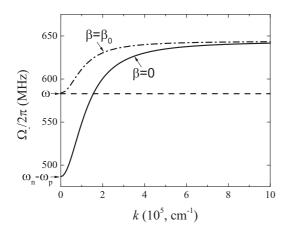


FIG. 8. The spectrum of *n*-magnons in MnCO₃ for zero deflection angle (solid line) and for the deflection angle given by Eq. (51) (dashed-dotted line). The frequency of rf pumping ω is indicated by a dashed line.

showed above, such a conventional heating scenario does not agree with the experiments described here.

In an alternative scenario, two *n*-magnons with wave vectors $\pm \mathbf{k}$ can mutually transform their $|\mathbf{k}|$ in momentum-conserving, four-magnon processes to the value $|\mathbf{k} - \delta \mathbf{k}|$, which corresponds to the resonance condition $\Omega_{-}(|\mathbf{k} - \delta \mathbf{k}|, \beta) = \omega$. Thus, the rf pumping and accumulation of magnons continue, leading to a further increase of the deflection angle β , and so on. The accumulation of magnons and increase of the deflection angle stops when the bottom of the frequency band of *n*-magnons reaches the rf pumping frequency,

$$\omega_n - \omega_p \cos \beta_0 = \omega \tag{51}$$

(see the dashed-dotted line in Fig. 8). In this scenario, all accumulated *n*-magnons, which provide the value of the deflection angle in the above equation, are transformed into *n*-magnons with k = 0. Such a macroscopic accumulation of magnons in a single k = 0 mode can be identified as a Bose-Einstein condensate of nuclear spin waves by analogy with the atomic BEC. In this state of the nuclear system, the vector of nuclear magnetization is deflected by a large angle β_0 from the equilibrium orientation (e.g., as large as 70° in the experiments described here), while the magnitude of the magnetization vector is conserved. This description agrees with the interpretation of experimental results observed earlier [7–9].

Investigation of the coupled electron-nuclear spin precession in easy-plane antiferromagnets is interesting in the context of conventional magnon BEC. Unlike the conventional atomic BEC obtained by cooling the atomic system, the magnon BEC is established by an external rf pumping that compensates for the loss of quasiparticles. The magnon BEC was first observed in the antiferromagnetic *B* phase of superfluid ³He [21,22]. Owing to its absolute purity and the specific form of the magnetic energy potential, which ensures the stability of BEC, this system became a test bed for the investigation of conventional magnon BECs with k = 0 showing close analogy to the atomic BEC [1,23]. Later, the stable state of magnon BEC was also obtained in the antiferromagnetic *A* phase of superfluid ³He immersed in aerogel [24,25]. Superfluid

³He-A is particularly interesting in the context of the work presented here. This system is a two-sublattice antiferromagnetic quantum liquid in which the magnetic part of the Hamiltonian exactly corresponds to that of the easy-plane solid antiferromagnets considered here [26]. For this reason, the latter systems were suggested for observation of conventional magnon BEC [27]. The significant advantage of these systems over superfluid ³He is that they require rather moderate cryogenic temperatures around 1 K for the observation and study of BEC properties and spin superfluidity. We should point out that the conventional magnon BEC with k = 0 discussed here has very different properties from those of e-magnon BEC observed in ferromagnetic yttrium iron garnet (YIG) films [28,29]. In the latter case, the minimum of the magnetic energy corresponds to spin waves with $k \neq 0$ and has some similarities with charged density waves in superconductors.

V. CONCLUSIONS

We present a description of coupled electron-nuclear spin systems using the coupled Landau-Lifshitz-Gilbert equations, which take into account an indirect relaxation of nuclear spins via the electron spin subsystem. In our theory, the magnitude of the nuclear magnetization is conserved for arbitrary large excitation powers, which is drastically different from the conventional heating scenario based on the phenomenological Bloch approach. In particular, we consider the coupled electron-nuclear spin motion in the easy-plane antiferromagnetic crystals with a weak magnetic anisotropy in the basal plain. The equations of motion are solved analytically, assuming small deviations of electron magnetization vectors from their equilibrium orientations, which allow us to linearize the corresponding equations of motion for the electron subsystem. The solutions obtained are compared with experimental nonlinear NMR signals obtained in a MnCO₃ sample at temperatures below 1 K, and good quantitative agreement is found. This suggests that at the high excitation rf powers used in the experiment, the nuclear magnetization vector can be deflected at very large angles, exceeding 70° . This result is drastically different from predictions of the standard heating scenario based on the phenomenological Bloch approach. In particular, the latter predicts a significant reduction of the magnitude of the nuclear magnetization vector under conditions of our experiment. The proposed theory brings together the properties of nonlinear NMR in magnetic systems in the solids considered here and the superfluid ³He, where the magnitude of the nuclear magnetization vector is also conserved. This provides the theoretical background for the study of BEC of nonequilibrium magnons in MnCO₃.

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APPENDIX: LINEARIZED EQUATIONS FOR ELECTRON SPIN PRECESSION

The system of coupled equations (3) and (4) contains 12 dynamical variables corresponding to three components for each of four magnetization vectors $\mathbf{M}_{1,2}$ and $\mathbf{m}_{1,2}$. This system conveniently splits into two independent subsystems, each consisting of six equations, if we introduce dynamical variables according to [14]

$$2M^{\alpha_{\pm}} = M_1^{\alpha_1} \pm M_2^{\alpha_2}, \quad 2m^{\alpha_{\pm}} = m_1^{\alpha_1} \pm m_2^{\alpha_2}, \qquad (A1)$$

where the upper indices are defined as $\alpha_{\pm} \in [x_{\pm}, y_{\pm}, z_{\pm}], \alpha_1 \in [x_1, y_1, z_1], \alpha_2 \in [x_2, y_2, z_2]$, and $M_i^{x_i}, M_i^{y_i}, M_i^{z_i}$ $(m_i^{x_i}, m_i^{y_i}, m_i^{z_i}), i = 1, 2$, are components of electron (nuclear) magnetization vectors in the corresponding reference frames (x_1, y_1, z_1) and (x_2, y_2, z_2) defined in Fig. 2. The components M^{α_-} correspond to the high-frequency branch of oscillations of the vectors $\mathbf{M}_{1,2}$, which is weakly affected by the interaction between electron and nuclear spins. For this reason, only the low-frequency branch of $\mathbf{M}_{1,2}$ is of interest for NMR. This branch is described by the components M^{α_+} and m^{α_+} . It is convenient to mark the indices x_+, y_+ , and z_+ , respectively, as ξ , η , and ζ . In what follows, we consider the static magnetic fields $H > H_c$. The field H_c is defined by the relation

$$H_c(H_c + H_D) = H_E H_n m_0 / M_0, \quad H_n = A M_0,$$
 (A2)

where M_0 and m_0 are the magnitudes of vectors $\mathbf{M}_1, \mathbf{M}_2$ and $\mathbf{m}_1, \mathbf{m}_2$, respectively. The inequality $H > H_c$ is responsible for the conditions

$$M^{\xi,\eta} \ll M_0, \quad M^{\zeta} \approx M_0.$$
 (A3)

In this approximation, the linearized equations of motion for M^{ξ} and M^{η} become [14]

$$\frac{1}{\gamma_e} \frac{dM^{\xi}}{dt} = (H_E + H_A)M^{\eta} - H_n m^{\eta} - \frac{1}{\gamma_e T_{2e}} M^{\xi},$$

$$\frac{1}{\gamma_e} \frac{dM^{\eta}}{dt} = -\frac{H(H + H_D)}{H_E} M^{\xi}$$

$$+ \frac{H + H_D}{H_E} M_0 h(t) + H_n m^{\xi} - \frac{1}{\gamma_e T_{2e}} M^{\eta},$$
 (A4)

where T_{2e} is the transverse relaxation time of the electron magnetization. We will use the applied rf magnetic field $\mathbf{h} \parallel \mathbf{z}$ in the form

$$h(t) = 2H_1 \cos \omega t. \tag{A5}$$

Note that the approximation (A3) used to obtain the linearized Eqs. (A4) corresponds to the following condition,

$$|\omega - \omega_e|T_{2e} \gg 1, \tag{A6}$$

where $\omega_e = \gamma_e [H(H + H_D)]^{1/2}$ is the frequency of uniform precession of vector **M** neglecting the interaction between electron and nuclear spins. The condition (A6) is well satisfied for typical rf frequencies close to the NMR frequency.

Equation (A4) can be simplified further. Noting that, since $M^{\eta} \gg m^{\eta}$ and $\gamma_e(H_E + H_A)T_{2e} \gg 1$, we can neglect the last two terms in the right-hand side of the first equation

in (A4). Moreover, since typically $H_E \sim 10^5 - 10^7$ Oe and $H_A \sim 10^2 - 10^4$ Oe, we can neglect H_A comparing with H_E . Using these approximations, we obtain from (A4),

$$\frac{d^2 M^{\xi}}{dt^2} + \frac{2}{T_{2e}} \frac{dM^{\xi}}{dt} + \omega_e^2 M^{\xi} = 2\gamma_e^2 (H + H_D) M_0 H_1 \cos \omega t + \gamma_e^2 H_n H_E m^{\xi}.$$
(A7)

The above equation is solved by using the Fourier transformations

$$M_{\Omega}^{\xi} = \int dt \, M^{\xi}(t) e^{i\,\Omega t}, \quad m_{\Omega}^{\xi} = \int dt \, m^{\xi}(t) e^{i\,\Omega t}, \quad (A8)$$

for which we obtain

$$M_{\Omega}^{\xi} \approx M_{\Omega}^{\xi}(H_1) + M_{\Omega}^{\xi}(m_{\Omega}^{\xi}), \tag{A9}$$

$$M_{\Omega}^{\xi}(H_1) = \frac{M_0}{\omega_n} \omega_1 [\delta(\Omega + \omega) + \delta(\Omega - \omega)], \quad (A10)$$

$$M_{\Omega}^{\xi}\left(m_{\Omega}^{\xi}\right) = \frac{2M_{0}}{\omega_{n}} \left(\omega_{p} - i\frac{\Omega}{T_{2n}\omega_{n}}\right) \frac{m_{\Omega}^{\xi}}{m_{0}}, \qquad (A11)$$

where for the sake of convenience we use the following definitions,

$$\omega_1 = \gamma_n \eta H_1, \tag{A12}$$

$$\eta = \frac{H_n}{H} \left(1 - \frac{\Omega^2}{\omega_e^2} \right)^{-1}, \tag{A13}$$

$$\omega_{p} = \frac{\omega_{n}}{2} \frac{H_{n}H_{E}}{H(H+H_{D})} \frac{m_{0}}{M_{0}} \left(1 - \frac{\Omega^{2}}{\omega_{e}^{2}}\right)^{-1}, \quad (A14)$$

$$\frac{1}{T_{2n}} = \frac{2\omega_n \omega_p}{T_{2e} \omega_e^2} \left(1 - \frac{\Omega^2}{\omega_e^2}\right)^{-1}.$$
 (A15)

To obtain the above equations, we used the expansion in the small parameter $\Omega/T_{2e} \ll (\omega_e^2 - \Omega^2)$. The influence of the relaxation is described by the imaginary part of M_{Ω}^{ξ} in (A11). Note that the above equations satisfy the condition $\operatorname{Im}(M_{\Omega=0}^{\xi}) = 0$, which means the absence of dissipation at $\Omega = 0$. It is important to fulfill this strict requirement imposed by the theory of irreversible processes.

The equations for the time-dependent components $m^{\xi}(t)$, $m^{\eta}(t)$, and $m^{\zeta}(t)$ of the nuclear magnetization vector can be obtained from Eqs. (4) if we exclude the corresponding timedependent components of the electron magnetization vector. This can be done using the obtained relation between the Fourier components M_{Ω}^{ξ} and m_{Ω}^{ξ} given by (A8)–(A11). The resulting integrodifferential equations are rather complicated and can be significantly simplified using the following approximations. First, we neglect the relaxation terms in Eqs. (4). This approximation is justified by the extremely large value of the hyperfine field $\mathbf{H}_{m_{1,2}}$ in the systems considered here. This approximation, in the spirit of the Landau-Lifshitz-Gilbert approach, is responsible for the conservation of the magnitude of the nuclear magnetization vector $m_0^2 = m^{\xi}(t)^2 + m^{\eta}(t)^2 +$ $m^{\zeta}(t)^2$. Second, we neglect the dependence of η , ω_p , and T_{2n} on Ω given by (A13)–(A15). This is justified for typical frequencies of nuclear spin precession, which are much smaller than the frequency of electron spin precession ω_e . Finally, we neglect harmonics terms at frequencies $2\omega, 3\omega, \ldots$ in equations for the time-dependent components $m_X(t)$, $m_Y(t)$, and $m_Z(t)$ defined by Eqs. (15). These approximations result in Eqs. (19)–(21), the solutions of which are discussed in the text.

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