Long-range interactions in antiferromagnetic quantum spin chains

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We study the role of long-range dipolar interactions on antiferromagnetic spin chains, from the classical $S \rightarrow \infty$ limit to the deep quantum case S = 1/2, including a transverse magnetic field. To this end, we combine different techniques such as classical energy minima, classical Monte Carlo, linear spin waves, bosonization, and density matrix renormalization group (DMRG). We find a phase transition from the already reported dipolar ferromagnetic region to an antiferromagnetic region for high enough antiferromagnetic exchange. Thermal and quantum fluctuations destabilize the classical order before reaching magnetic saturation in both phases, and also close to zero field in the antiferromagnetic phase. In the extreme quantum limit S = 1/2, extensive DMRG computations show that the main phases remain present with transition lines to saturation significatively shifted to lower fields, in agreement with the bosonization analysis. The overall picture maintains a close analogy with the phase diagram of the anisotropic XXZ spin chain in a transverse field.

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I. INTRODUCTION

Long-range interactions in quantum systems have recently attracted much attention. While short-range interactions are naturally present in quantum gases, longer-range interactions are much harder to control. To investigate them, ultracold gases of particles with large magnetic or electric dipole moments, atoms in Rydberg states, or cavity-mediated interactions have been studied [1]. These experiments also open the possibility of simulating dipole-dipole interactions in one-dimensional spin chains [2]. Since then, theoretical and numerical investigations of dipolar spin chains have been revitalized [3]. On the other hand the inclusion of dipolar and more generally long-ranged interactions in classical and quantum models has proven to modify in different degrees the outcoming physics [4–6].

Motivated by these studies, we consider the competition between short-range antiferromagnetic exchange and longrange dipolar interactions in spin chains in order to explore the presence of novel phases, either ordered or disordered. We also include an external magnetic field, transverse to the dipole-dipole induced anisotropy, which competes with both antiferromagnetic and dipolar classical ordering.

The present analysis follows different approaches that allow us to explore from the classical $S \rightarrow \infty$ limit to the deep quantum case S = 1/2, as well as parameter regions where antiferromagnetic exchange dominates over dipolar interactions and the other way around. For large *S* we perform classical energy analysis, classical Monte Carlo simulations, and linear spin wave fluctuations. On the other extreme, for spin S = 1/2 we employ bosonization techniques and density matrix renormalization group (DMRG) computations.

With these techniques, we first draw a semiclassical phase diagram showing three distinctive phases: a dipole-induced ferromagnetic phase, a disordered Luttinger liquid phase, and an antiferromagnetic Ising Néel phase. The stability of the boundaries between those phases and their transition to the magnetically saturated phase is discussed in terms of thermal and quantum spin wave fluctuations. Beyond fluctuations, for the S = 1/2 deep quantum case our main results show a substantial reduction of the magnetic field rendering both the dipolar and antiferromagnetic ordered phases into the

magnetically saturated phase. The quantum phases found are in correspondence with those present in an anisotropic S = 1/2 spin chain; in this sense the main effect of dipolar interactions seems to be encoded in their nearest-neighbor contribution.

The paper is organized as follows. In Sec. II we define the model and review the classical magnetic phases. In Sec. III we discuss the effect of thermal and quantum fluctuations by means of Monte Carlo simulations and spin wave calculations. Our main results for spin S = 1/2 are presented in Sec. IV through extensive DMRG data, preceded by a bosonization analysis. We close this work with concluding remarks in Sec. V.

II. MODEL AND CLASSICAL DESCRIPTION

We consider a spin chain in the x direction with long-range dipolar interactions and nearest-neighbor antiferromagnetic exchange coupling J > 0 (see Fig. 1). The Hamiltonian reads

$$\mathcal{H} = J \sum_{i} \vec{S}_{i} \cdot \vec{S}_{i+1} + \mu^{2} \sum_{i < j} \left(\frac{\vec{S}_{i} \cdot \vec{S}_{j}}{|\vec{r}_{i} - \vec{r}_{j}|^{3}} - 3 \frac{S_{i}^{x} S_{j}^{x}}{|\vec{r}_{i} - \vec{r}_{j}|^{3}} \right) - h \sum_{i} S_{i}^{z}, \tag{1}$$

where \vec{S}_i is a dimensionless spin *S* operator at site $\vec{r}_i = i a \vec{x}$ with *a* the lattice spacing between consecutive spins and μ the effective gyromagnetic moment of the spins. Since $|\vec{r}_i - \vec{r}_j| = a|i - j|$ we will use in the following a dipolar coupling $D = \frac{\mu^2}{a^3}$ with units of energy. An external magnetic field $\vec{h} = h\vec{z}$ is chosen to be perpendicular to the anisotropy introduced by dipolar interactions along the spin chain direction.

In this section we review the classical ground configurations of the Hamiltonian in Eq. (1). Dipolar interactions tend to align classical spins along the x axis while the transverse magnetic field induces a tilting towards the z axis. Ferromagnetic exchange couplings J < 0 are fully satisfied in such configurations [3], but the present antiferromagnetic couplings J > 0 would prefer to set a Néel order in the xy plane. One can then search for the classical ground state configurations



FIG. 1. Schematic representation of nearest-neighbor exchange interactions and long-range dipolar interactions. Transverse magnetic field is also depicted.

within the manifold parametrized by

$$\vec{S}_i = S(\sin\theta\cos\phi, (-1)^i\sin\theta\sin\phi, \cos\theta), \qquad (2)$$

where θ is the tilting angle with respect to the magnetic field and ϕ describes a staggered deviation from the chain direction into the *xy* plane. The energy per site of such configurations reads

$$\frac{E_0}{S^2} = J \left[\cos^2 \theta + \sin^2 \theta \cos(2\phi) \right] + \zeta(3) D \left(\cos^2 \theta - 2\sin^2 \theta \cos^2 \phi - \frac{3}{4}\sin^2 \theta \sin^2 \phi \right) - \frac{h}{S}\cos \theta, \qquad (3)$$

where $\zeta(3) = \sum_{n=1}^{\infty} \frac{1}{n^3} \approx 1.20206$ stems from the numerical evaluation of the Riemann zeta function.

It is readily found that this classical energy, for $J \leq J_c \equiv \frac{5}{8}\zeta(3)D$ and below a saturation field $h_{\text{sat}}^{\text{DF}}(D) = 6\zeta(3)SD$, exhibits a minimum at $\phi = 0$ (parallel spins in the *xz* plane) and tilting angle

$$\theta^{\rm DF}(h,D) = \arccos\left(\frac{h}{6\zeta(3)SD}\right)$$
(4)

defining a "dipolar-ferromagnetic" (DF) phase. This phase has a \mathbb{Z}_2 mirror degeneracy under exchange $\theta \to -\theta$. Instead, for $J \ge J_c$ and below a saturation field $h_{\text{sat}}^{\text{AF}}(J,D) = 4S[J + \frac{7}{2}\zeta(3)D]$, the minimum appears at $\phi = \pm \pi/2$ and

$$\theta^{\rm AF}(h,J,D) = \arccos\left(\frac{h}{4S(J + \frac{7}{8}\zeta(3)D)}\right) \tag{5}$$

defining an antiferromagnetic (AF) phase where spins lie in staggered tilted directions in the yz plane. This phase has a \mathbb{Z}_2 discrete translation degeneracy along the chain direction. The classical phase diagram in the *h* vs *J* plane is shown in Fig. 2.

At the specific value $J = J_c > 0$ the ground state manifold is infinitely degenerate for any value of the external magnetic field below saturation, with a free value for the angle ϕ that can continuously interpolate between the dipolar-ferromagnetic and the antiferromagnetic phases. Also notice that at zero field the classical AF phase possesses an extra U(1) degeneracy associated with rotations around the chain direction. Such degeneracies enhance the role of classical and quantum fluctuations, as we discuss in the next section.



FIG. 2. Classical phases are described in the *h* vs *J* plane: tilted dipolar ferromagnetic (DF), tilted antiferromagnetic (AF), and magnetic saturation (FM). Shaded regions indicate predominance of both thermal and spin wave quantum fluctuations. We show below that the boundaries to saturation are significantly shifted down in the quantum S = 1/2 limit.

III. CLASSICAL AND QUANTUM FLUCTUATIONS

A. Monte Carlo simulations

We study here the effects of classical thermal fluctuations on the model presented in Sec. II, in order to check the stability of the phase boundaries of the zero-temperature classical phase diagram shown in Fig. 2. To this end we have run Monte Carlo simulations using the standard Metropolis algorithm combined with overrelaxation (microcanonical) updates for chains of L = 300 sites. Finite-size systems with periodic boundary conditions are considered, so that long-range interactions are taken up to L/2 neighbors. For each simulation 10^4 Monte Carlo steps (mcs) were dedicated to thermalization, lowering the temperature with the annealing technique at a rate $T_{n+1} = 0.9 \times T_n$. Measurements are then taken during the 2×10^4 subsequent mcs. The results presented for each data point describe the average over 100 independent simulations.

In order to describe the effects of thermal fluctuations in both DF phase, $J < J_c$, and AF phase, $J > J_c$, we define two different order parameters: a mean magnetization *m* defined as

$$m = \sqrt{m_x^2 + m_y^2 + m_z^2},$$
 (6)

where $m_{\alpha} = \frac{1}{L} \sum_{i} S_{i}^{\alpha}$ ($\alpha = x, y, z$), and an antiferromagnetic magnetization m^{AF} defined as

$$m^{\rm AF} = \sqrt{m_x^2 + (m_y^{\rm stag})^2 + m_z^2},$$
 (7)

where $m_y^{\text{stag}} = \frac{1}{L} \sum_i (-1)^i S_i^y$ picks the staggered contribution of the *y* components.

We show *m* and m^{AF} as functions of the external magnetic field for different values of *J* and T/D in Fig. 3. The top panel shows *m* for two values of $J < J_c$ at T/D = 0.02. The external magnetic field is normalized by the corresponding T = 0 saturation value h_{sat}^{DF} (see Sec. II). There is a clear dip at $h/h_{sat}^{DF} < 1$ for all values of *J*, due to temperature effects. This is illustrated in the inset of the top panel, which shows *m* vs h/h_{sat}^{DF} for J/D = 0.5 at different temperatures. Analogously,



FIG. 3. Order parameters *m* (top) and m^{AF} (bottom) as functions of the external magnetic field scaled with the corresponding T = 0 saturation values. Temperature is set at T/D = 0.02 for different values of J/D for $J < J_c$ (top) and $J > J_c$ (bottom).

the bottom panel shows m^{AF} versus h/h_{sat}^{AF} for three values of $J > J_c$ and T/D = 0.02, while the inset shows the effects of temperature for J/D = 1.

A dip close to saturation is observed, as expected, in both DF and AF phases which increases with temperature. In the AF phase, on the other hand, one observes a very pronounced dip in the staggered magnetization close to zero field due to the U(1) symmetry of the AF classical solution in absence of magnetic field.

B. Spin wave spectrum

Linear spin wave (LSW) fluctuations around the classical $S \rightarrow \infty$ solutions can be analyzed in a standard way. One first introduces local axes such that at each site *i* a new *z'* axis coincides with the classical solution spin orientation. Within the LSW approximation, the spin operator components in the local axes can be represented by bosonic Holstein-Primakoff local operators a_i , a_i^{\dagger} as

$$S_{i}^{z'} = S - a_{i}^{\dagger} a_{i},$$

$$S_{i}^{x'} = \sqrt{\frac{S}{2}} (a_{i}^{\dagger} + a_{i}),$$

$$S_{i}^{y'} = i \sqrt{\frac{S}{2}} (a_{i}^{\dagger} - a_{i}).$$
(8)

After a Fourier transformation in a periodic chain with L sites,

$$a_i = \frac{1}{\sqrt{L}} \sum_k e^{ikx_i} a_k, \tag{9}$$

where $k = p \frac{2\pi}{La}$, p = -L/2, ..., L/2, and $x_i = i a$, and ignoring cubic and higher order terms in a_k, a_k^{\dagger} , one can get the

form

$$\mathcal{H} = S \sum_{k} \{ A_{k} (a_{k}^{\dagger} a_{k} + a_{-k}^{\dagger} a_{-k}) + B_{k} (a_{k}^{\dagger} a_{-k}^{\dagger} + a_{k} a_{-k}) \}$$
(10)

with real $A_k \ge |B_k|$. Then the Hamiltonian can be diagonalized by a Bogoliubov transformation

$$a_{k} = u_{k}\alpha_{k} + v_{k}\alpha_{-k}^{\dagger},$$

$$a_{-k}^{\dagger} = v_{k}^{*}\alpha_{k} + u_{k}^{*}\alpha_{-k}^{\dagger},$$
(11)

with $|u_k|^2 - |v_k|^2 = 1$ ensuring that $\alpha_k, \alpha_k^{\dagger}$ are bosonic modes. The Hamiltonian finally reads

$$\mathcal{H} = NE_0 + \frac{S}{2} \sum_{k} (\varepsilon_k - A_k) + S \sum_{k} \varepsilon_k \alpha_k^{\dagger} \alpha_k, \qquad (12)$$

where E_0 is the classical energy per site given in Eq. (3) and $\varepsilon_k = \sqrt{A_k^2 - |B_k|^2}$ are the Bogoliubov mode energies.

At the dipolar-ferromagnetic phase an appropriate global orthogonal coordinate system is set by rotating the original axes an angle $\theta_{\text{DF}}(h, D)$ around the *y* direction. The Hamiltonian coefficients A_k^{DF} , B_k^{DF} in Eq. (10) are given by

$$A_{k}^{\mathrm{DF}} = J[\cos(ka) - 1] + D\left(3\sin^{2}\theta^{\mathrm{DF}} - 1\right)\left[\zeta(3) + \frac{1}{2}F(ka)\right] + \frac{h}{2S}\cos\theta^{\mathrm{DF}},$$
$$B_{k}^{\mathrm{DF}} = -\frac{3}{2}D\cos^{2}\theta^{\mathrm{DF}}F(ka), \qquad (13)$$

where $F(ka) \equiv \text{Re}[\text{Li}_3(e^{ika})]$ with $\text{Li}_3(z)$ the polylogarithm¹ series $\text{Li}_3(z) \equiv \sum_{n=1}^{\infty} \frac{1}{n^3} z^n$.

At the antiferromagnetic phase the appropriate axes are local ones, obtained by rotating the original axes at each site *i* with an angle $(-1)^i \theta^{AF}(h, J, D)$ about the *x* direction. The Hamiltonian coefficients A_k^{AF} , B_k^{AF} in Eq. (10) are then given by

$$A_{k}^{AF} = J\cos^{2}\theta^{AF}\cos(ka) - J\cos(2\theta^{AF}) + D\left(\frac{1}{2}\cos^{2}\theta^{AF} - 1\right)F(ka) + \frac{1}{2}D\sin^{2}\theta^{AF}G(ka) - D\zeta(3)\left(\cos^{2}\theta^{AF} - \frac{3}{4}\sin^{2}\theta^{AF}\right) + \frac{h}{2S}\cos\theta^{AF}, B_{k}^{AF} = J\sin^{2}\theta^{AF}\cos(ka) - D\left(\frac{1}{2}\cos^{2}\theta^{AF} + 1\right)F(ka) - \frac{1}{2}D\sin^{2}\theta^{AF}G(ka),$$
(14)

where $G(ka) \equiv \frac{1}{4}F(2ka) - F(ka)$.

In either case, the semiclassical ground state is the Bogoliubov vacuum $|0\rangle$ annihilated by the operators α_k . The ground state energy is then given by

$$\mathcal{H} = NE_0 + \frac{S}{2} \sum_k (\varepsilon_k - A_k).$$
(15)

Within the LSW framework the sensible order parameter to compute is the average of the local magnetizations along the

 $^{{}^{1}}F(ka)$ is also known as the Clausen function $Cl_{3}(ka)$.



FIG. 4. Mean local magnetization vs h for different values of antiferromagnetic $J < J_c$, in the LSW approximation.

classical directions z', defined as

$$m^{z'} = \frac{1}{N} \sum_{i} \langle 0|S_i^{z'}|0\rangle = S + \frac{1}{2} - \sum_{k} \frac{A_k}{2\sqrt{A_k^2 - |B_k|^2}}.$$
 (16)

A large value of the summation in the last term signals the breakdown of the LSW approximation, meaning that quantum fluctuations destroy the classical order. This occurs in the shaded regions of the classical phase diagram in Fig. 2.

We show in Fig. 4 $m^{z'}$ vs *h* for coupling ratios within the DF phase, $J/D < \frac{5}{8}\zeta(3)$, while in Fig. 5 we show the corresponding results for ratios within the AF phase, $J/D > \frac{5}{8}\zeta(3)$.

These results extend those obtained in Ref. [3], at J = 0, to the whole DF region and show a new phase with antiferromagnetic characteristics. One can observe that the LSW corrections to the order parameter diverge as h approaches h_{sat} , both in the DF and the AF phases. From the slopes in these figures the sensitivity to quantum fluctuations is shown to be higher when J approaches J_c , i.e., when the competition between dipolar and exchange interactions is stronger. In the AF phase quantum fluctuations are crucial not only close to saturation but also close to zero field, as expected for antiferromagnetic systems. Here the LSW corrections are more important for larger J. Indeed, the crossings in the curves in Fig. 5 signal a crossover from a dipolar-reminiscent behavior, close to J_c , towards an exchange-dominated behavior for $J \gg D$.

These features agree with the classical thermal fluctuations picture and are confirmed by extensive DMRG computations and a bosonization analysis in the next section.



FIG. 5. Mean local magnetization vs *h* for different values of antiferromagnetic $J > J_c$, in the LSW approximation.

(17)

IV. EXTREME QUANTUM LIMIT: SPIN-1/2 CASE

In this section we discuss specific features of the spin-1/2 case, using bosonization techniques and extensive state-of-theart DMRG numerical computations for long-range coupled systems.

A. Bosonization approach

For the present discussion we find it convenient to rewrite the Hamiltonian in Eq. (1) by separating the nearest-neighbor interactions (\mathcal{H}_{NN}) from the longer-distance dipolar terms (\mathcal{H}_{int}), which we then treat perturbatively. For the sake of clarity, we consider first the zero-field case. The Hamiltonian reads

 $\mathcal{H} = \mathcal{H}_{\rm NN} + \mathcal{H}_{\rm int},$

where

$$\mathcal{H} = (J+D)\sum_{i} \left[S_{i}^{y} S_{i+1}^{y} + S_{i}^{z} S_{i+1}^{z} + \frac{J-2D}{J+D} S_{i}^{x} S_{i+1}^{x} \right]$$
(18)

and

$$\mathcal{H}_{\text{int}} = D \sum_{j>i+1} \left(\frac{-2S_i^x S_j^x + S_i^y S_j^y + S_i^z S_j^z}{(j-i)^3} \right).$$
(19)

Notice that \mathcal{H}_{NN} corresponds to the well known anisotropic XXZ Heisenberg chain with $\Delta = \frac{J-2D}{J+D}$ (see for instance Ref. [7]). In this sense, the short-range effect of dipolar interactions is the onset of an exchange-like anisotropy along the chain direction. Without further interactions, a Luttinger liquid phase is present, extending from a ferromagnetic transition point at J = 0.5D ($\Delta = -1$) up to an isotropic Heisenberg point ($\Delta = 1$) to be reached at $J/D \rightarrow \infty$, with a free fermion point ($\Delta = 0$) located at J = 2D. Bosonization of the spin S = 1/2 chain in this regime has succeeded in describing the ground state and correlation functions, as well as allowing for a conformal perturbative scheme (see for instance Ref. [8]). Following this scheme one can show that the $1/r^3$ decaying long-range dipolar interactions in \mathcal{H}_{int} do not alter the Luttinger liquid behavior, but only renormalize the Luttinger parameters. In accordance with the spin wave indications, the system enters a disordered phase for $J > J_c^q$, a quantum critical point which is eventually shifted from 0.5D by dipolar interactions.

In the region J < 0.5D the truncated Hamiltonian \mathcal{H}_{NN} enters a gapped ferromagnetic phase [7], with a twofold degenerate ground state characterized by the order parameter $\langle S_i^x \rangle \neq 0$. Though conformal perturbations cannot be applied in this region, notice that the full Hamiltonian $\mathcal{H}_{NN} + \mathcal{H}_{int}$ classically exhibits the same ordering, which is robust against thermal and quantum fluctuations. For the quantum case S = 1/2 we explore numerically this order parameter in the next section.

The behavior of the present system under a transverse magnetic field, both in the Luttinger and the ferromagnetic region, can be naturally related to the anisotropic XXZ Heisenberg chain in a transverse field. As discussed in Refs. [9] and [10], a small transverse field induces a gap on the Luttinger



FIG. 6. Total magnetic moment m/S as a function of the applied magnetic field *h* for two representative values of the antiferromagnetic exchange $J/D = 0.6 < J_c$ and $J/D = 1.0 > J_c$ and lattice sizes L = 36,48,64 (each curve is normalized by its corresponding saturation magnetic field h_{sat}^{DF} and h_{sat}^{AF}).

phase, driving the spin system into an Ising Néel phase with staggered nonvanishing expectation value of spin components along the *y* direction (i.e., transverse to both the anisotropy and the magnetic field). On the gapped ferromagnetic phase, the transverse field diminishes the order parameter, until a quantum phase transition into a paramagnetic phase is reached. Discussing how the longer-range interactions in \mathcal{H}_{int} may modify this picture is beyond the scope of the present paper.

These predictions, namely the renormalization of the quantum critical point and the behavior in a transverse magnetic field, are confirmed below by a DMRG analysis.

B. DMRG calculations

With the aim of characterizing the spin S = 1/2 quantum phases of the antiferromagnetic chain with dipolar interactions, we have analyzed the ground state of the Hamiltonian in Eq. (1) with the DMRG technique [11]. Including long-range interactions is a nontrivial task, comparable with current studies of two-dimensional spin systems [12,13]. Based on our experience in such investigations, we use here open boundary conditions and long-range interactions involving all available neighbors in chains of finite size. We have considered chains of length *L* up to 64 sites, keeping m = 350 states and achieving truncation errors in the density matrix of the order of 10^{-11} .

As a first step to identify the configurations of the system we compute the total magnetic moment m defined in Eq. (6), where now $m_{\alpha} = \frac{1}{L} \sum_{i} \langle S_{i}^{\alpha} \rangle$ stands for the average of quantum expectation values. In Fig. 6 we show DMRG results for the total magnetic moment m as a function of the magnetic field for two representative values J = 0.6D and J = 1.0Dof the antiferromagnetic exchange interaction, chosen to be compared with the figures shown in Sec. III. As in the classical chain, we observe two different configurations depending on the value of the exchange interaction J being below or above a critical value. In more detail (not shown in the figure) we have been able to estimate a quantum critical coupling $J_c^q =$ 0.7275(25) slightly lower than the classical value obtained in Sec. II, $J_c = \frac{5}{8}\zeta(3)D \approx 0.7512$. In the DF phase, $J < J_c^q$, the magnetization exhibits a minimum, a similar behavior to that of the pure dipolar case in Ref. [3]. Instead, in the AF phase, $J > J_c^q$, we find that the total magnetization smoothly increases with the applied magnetic field.



FIG. 7. Magnetic moments m_x and m_z as a function of the applied magnetic field for a chain of L = 48 sites. Left panel: Results for $J/D = 0.6 (J < J_c)$. Right panel: Results for $J/D = 1.0 (J > J_c)$.

As a further step we analyze the components of the magnetic moments along the three directions x, y, and z separately. In Fig. 7 we show the values of m_x and m_z as functions of the applied magnetic field, for exchange $J = 0.6D < J_c^q$ and J = $1.0D > J_c^q$. In both phases m_z increases smoothly with the applied field, approaching to saturation at about the classical saturation field. Instead, the m_x component makes apparent the difference between the DF and AF regimes. The component m_y vanishes in both cases, and for the sake of clarity it is not shown in the figures; we argue below that the reason why this happens is very different for each phase.

In the DF phase $(J < J_c^q)$ we found a twofold-degenerate ground state, as dictated by parity \mathbb{Z}_2 symmetry under $x \rightarrow z_2$ -x reflections. A parity-resolved basis for this ground state subspace is given by states with $m_x < 0$ and $m_x > 0$. A tiny magnetic field $h_x = 10^{-10} D$ acting just on the end sites of the chain [3] is enough to explicitly break parity \mathbb{Z}_2 symmetry, selecting the state with $m_x > 0$. This response supports the interpretation that the finite-size ground state is a simple superposition of disentangled ferromagnetic product states. Following this strategy we produced the states shown in the left panel of Fig. 7: at low fields the most important contribution to the magnetization is provided by the m_x component, followed by a sudden drop well before m_z approaches saturation. This explains the pit in Fig. 6. As the magnetic field increases, we found a magnetic field $h_c \approx 0.47 h_{\text{sat}}^{DF}$ above which all the magnetization weight is already in the z direction. The value of h_c is obtained by extrapolation of the $h_c(L)$ for different chain lengths, and signals that all spins already align with the magnetic field at less than half the classical saturation field. In this phase, as in the pure dipolar case, there is no symmetry reason for the system to choose an orientation in the y direction; we accordingly found that $\langle S_i^y \rangle = 0$ at each site.

In the AF phase $(J > J_c^q)$ we observed distinct ground states in the absence or presence of the external magnetic field. Without external field, in agreement with the Luttinger regime found in the bosonization analysis, the ground state shows no order; local expectation values vanish for any spin component as shown in the right panel of Fig. 7 at h = 0. Under a magnetic field, also in agreement with the gapped Ising Néel phase predicted by bosonization, we found a twofold-degenerate ground state, related to one-site \mathbb{Z}_2 translation invariance. While we expect a spontaneous



FIG. 8. End-to-end correlations for lattice lengths L = 36,48,64and the $L \rightarrow \infty$ extrapolation as a function of the applied magnetic field for J/D = 1.0.

symmetry breaking in the thermodynamic limit, leading to a staggered magnetization along the y direction, one should notice that translation symmetry is not broken in the available finite-size systems;² as a consequence, one-point operators can only show homogeneous expectation values and a staggered magnetization is compatible with the observed local result $\langle S_i^y \rangle = 0$. In contrast to the DF phase, in this case a tiny staggered magnetic field h_{y} acting on the end sites of the chain is not enough to explicitly break \mathbb{Z}_2 translation invariance. In order to obtain clear signals of the Néel order classically observed and quantum mechanically predicted in the AF phase, we resorted to the computation of S^{y} two-point spin correlations $\langle S_i^y S_j^y \rangle$ which allow for staggered nonvanishing results, invariant under one-site translations. For $J > J_c^q$ we indeed found staggered correlations, which include an exponentially decaying connected part and a nonvanishing long-range order disconnected part,

$$\left\langle S_{i}^{y}S_{j}^{y}\right\rangle = \left\langle S_{i}^{y}S_{j}^{y}\right\rangle_{0} + \left\langle S_{i}^{y}\right\rangle\left\langle S_{j}^{y}\right\rangle \tag{20}$$

with $\langle S_i^y \rangle \langle S_j^y \rangle \propto (-1)^{j-i} (m_y^{\text{stag}})^2$. We thus can extract the antiferromagnetic order parameter from the two-point spin correlations. The long-distance behavior can be better analyzed by considering the end-to-end correlations³

$$C_{1,L} = \left\langle S_1^y S_L^y \right\rangle. \tag{21}$$

As pointed out in Ref. [14] these correlations between the spins at opposite ends of the chain provide a more tractable distance scaling than bulk correlations. Because of the evenodd sign of correlators, we rather plot $|C_{1,L}|$ to identify the response of end-to-end correlations to the applied magnetic field. We plot in Fig. 8 the results for the considered finite-size chains at J/D = 1.0 for several values of the field, as well as their infinite-size extrapolations. As a first feature it can be observed that at intermediate values of the field, the order parameter is nonzero, with a negligible size scaling. This confirms the presence of long-range AF order for $J > J_c^q$. As a second one, the staggered correlations drop down and vanish at about 60% of the classical saturation field. This coincides with the high slope rise of m_z in the right panel of Fig. 7. Finally, end-to-end correlations approach zero as the magnetic field vanishes (this is more clear deep inside the AF phase, for instance at J/D = 1.5), confirming that staggered order is destroyed at zero field as suggested by thermal and quantum fluctuations and symmetry arguments. Moreover, the noticeable size scaling indicates the typical quasi-long-range order in the Luttinger phase. These results resemble those found for the Luttinger phase of the anisotropic XXZ spin chain in the presence of a transverse field (see for instance Fig. 5 in Ref. [10]).

V. CONCLUSIONS

In the present work we have described the phase diagram for an antiferromagnetic nearest-neighbor spin chain (with exchange strength J) including the effects of long-range dipole-dipole interactions (of strength D) and a transverse magnetic field (of strength h).

On the one hand, we have characterized the presence of ordered phases for classical spins and their stability under the influence of classical and quantum fluctuations. The main feature here is the presence of a phase transition at $J/D = \frac{5}{8}\zeta(3)$ from a dipolar-dominated phase to an antiferromagnetic phase. In the former one classical spins are aligned in a ferromagnetic pattern forming an angle $\theta^{\text{DF}}(h,D) = \arccos(\frac{h}{6\zeta(3)SD})$ with the external field, according to the competition between the dipolar tendency to align them parallel to the chain and the Zeeman energy, while in the latter phase spins order antiferromagnetically, transverse to the dipolar anisotropy and canted towards the external field at an angle $\theta^{\text{AF}}(h, J, D) = \arccos(\frac{h}{4S[J+\frac{2}{8}\zeta(3)D]})$. Classical and quantum fluctuations destroy both these classical orders before reaching the saturation field, and also the antiferromagnetic order close to zero field.

On the other hand, in the extreme quantum case S = 1/2, DMRG computations indicate that classical order disappears well before reaching the classical saturation fields. This is in agreement with previous studies on the purely dipolar S = 1/2chain [3] where a quantum critical point belonging to the 2dIsing universality class was identified and the effect of quantum fluctuations was proved to reduce the value of the critical field to magnetic saturation. Regarding the competition between dipolar ferromagnetic and antiferromagnetic orders, we have found that the quantum critical point separating these phases is slightly shifted (about 3%) to a lower value. This behavior is fully compatible with the bosonization analysis discussed in the text. The observed quantum phases are similar to those present in the well known XXZ anisotropic spin chain. Also the quantum critical transition reported in Ref. [3] for the dipolar S = 1/2 chain and the corresponding transition line for the XXZ chain belong to the same universality class. Our results suggest that the qualitative effect of dipolar interactions might be encoded in the anisotropy introduced by the nearestneighbor terms. Were this the case, the shaded phase transition lines in Fig. 2 would be critical. Further work along this rationale is in progress and will be reported elsewhere.

²The argument is valid for periodic boundary conditions. Still, onesite translations are an approximate symmetry for the open boundary chains considered in this work.

³In order to reduce boundary effects caused by OBC we discarded some sites at each end of the chain.

The current development of quantum simulations with ultracold trapped atoms is promising for testing an ever wider spectrum of quantum many-body properties. Some key achievements paving the way to the present discussion are the techniques described in Ref. [15] to simulate spin-1/2 exchange interactions and the control of magnetic dipoledipole interactions between chromium atoms in Ref. [16], as well as the stronger electric dipole-dipole interactions between polar molecules in Ref. [17]. Most recently, a controlled competition between short-range and long-range interactions in a bosonic optical lattice revealed in Ref. [1] the possibility of simulating different quantum phases and their transitions.

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We hope that such techniques can test our predictions in the near future.

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