The spin-$\frac{1}{2}$ Heisenberg ferromagnet on the pyrochlore lattice: A Green’s function study

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We consider the pyrochlore-lattice quantum Heisenberg ferromagnet and discuss the properties of this spin model at arbitrary temperatures. To this end, we use the Green’s function technique within the random-phase (or Tyablikov) approximation as well as the linear spin-wave theory and quantum Monte Carlo simulations. We compare our results to the ones obtained recently by other methods to corroborate our findings. Finally, we contrast our results with the ones for the simple-cubic-lattice case: both lattices are identical at the mean-field level. We demonstrate that thermal fluctuations are more efficient in the pyrochlore case (finite-temperature frustration effects). Our results may be of use for interpreting experimental data for ferromagnetic pyrochlore materials.

Key words: quantum spin system, Heisenberg model, pyrochlore lattice, Green’s functions, random-phase approximation

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The Green’s function method in quantum many-body theory emerged in the second half of the twentieth century. At that time, several outstanding books and reviews such as the ones by Kadanoff and Baym [1], Abrikosov, Gorkov, and Dzyaloshinski [2], Zubarev [3], Bonch-Bruevich and Tyablikov [4], Elk and Gasser [5] etc. appeared. Professor I.V. Stasyuk was among the pioneers in Ukraine who began to use these new methods of quantum statistics in solid state researches. His Green’s function papers on ferroelectric and magnetic crystals, molecular excitons or on diagram technique for the Hubbard model written in the late sixties–early seventies are well known in the community [6–10]. Moreover, he was lecturing for graduates and post-graduates for many years. Later on he wrote an excellent textbook on Green’s functions [11] where one can find many details of calculations as well as various applications in solid state physics.

In what follows we consider one specific quantum spin-lattice system — the spin-$\frac{1}{2}$ Heisenberg pyrochlore ferromagnet — using the Green’s function formalism. Clearly, for about sixty years a lot of results were obtained for various quantum spin-lattice systems. However, the Green’s function technique, even with utilizing a simple random-phase (or Tyablikov) approximation, is still used in recent studies for exploring complicated spin-lattice models (see, e.g., [12–14]). Moreover, while writing this paper we also bear in mind a pedagogical goal intending to explain all calculations and compare our findings with the known outcomes of other approaches. Last but not least, the model we discuss is related to frustrated quantum magnets — a hot topic during last thirty years.

It is our pleasure to dedicate this paper to the upcoming eighties birthday of Professor I.V. Stasyuk, who has influenced many studies in Lviv, including our ones, over about last fifty years.
1. Introduction

One of the standard three-dimensional models for a study of geometrically frustrated spin systems is the pyrochlore Heisenberg antiferromagnet. It is known that the nearest-neighbor exchange interactions, because of the lattice geometry (a three-dimensional network of corner-sharing tetrahedra), cannot establish magnetic ordering even at zero temperature \( T = 0 \). Besides the purely theoretical interest in the spin-system physics of this celebrated model, there are various real materials which can be described with the help of the antiferromagnetic Heisenberg model on the pyrochlore lattice.

Less attention has been paid to the pyrochlore-lattice Heisenberg ferromagnet. Clearly, the set of the eigenvalues of the Heisenberg Hamiltonian (energy levels) does not depend on the sign of the exchange interaction. As a result, the low-energy levels for the pyrochlore-lattice eigenvalues of the Heisenberg Hamiltonian (energy levels) does not depend on the sign of the exchange interaction. Therefore, the set of the eigenvalues of the Heisenberg Hamiltonian (energy levels) does not depend on the sign of the exchange interaction. As a result, the low-energy levels for the pyrochlore-lattice ferromagnet are the high-energy levels for the pyrochlore-lattice Heisenberg ferromagnet and they may manifest themselves only in finite-temperature thermodynamics, when due to thermal fluctuations they can come into play. Several systematic theoretical studies demonstrated that this is really the case. For example, it was found that the Curie temperature for the pyrochlore lattice is lower than for the simple-cubic lattice, although both lattices have the same number of nearest neighbors and thus are identical at the mean-field level.

From the experimental side, there is a number of compounds, the magnetic properties of which can be discussed by the same approach.

In what follows, we first introduce the model and present the corresponding linear spin-wave theory (section 2). In section 3, we describe the Green’s function approach to the model and in section 4, we discuss the low- and intermediate-temperature properties of the model making connections to previous studies. In section 5, we close the paper with a short summary pointing out some related problems which can be discussed by the same approach.

2. The model and the linear spin-wave theory

We begin with a brief description of the model. The three-dimensional pyrochlore lattice can be viewed as four interpenetrating face-centered-cubic sublattices. We set the edge length of the cubic cell of each face-centered-cubic sublattice to unity. Furthermore, we denote the origins of these face-centered-cubic sublattices as \( \mathbf{r}_1 = (0, 0, 0) \), \( \mathbf{r}_2 = (0, \frac{1}{2}, \frac{1}{2}) \), \( \mathbf{r}_3 = (\frac{1}{2}, 0, \frac{1}{2}) \), and \( \mathbf{r}_4 = (\frac{1}{2}, \frac{1}{2}, 0) \). For each face-centered-cubic sublattice, the position of its lattice sites is given by three integers \( m_1 \), \( m_2 \), and \( m_3 \), i.e., by \( \mathbf{R}_m = m_1 \mathbf{e}_1 + m_2 \mathbf{e}_2 + m_3 \mathbf{e}_3 \) with \( \mathbf{e}_1 = (0, \frac{1}{2}, \frac{1}{2}) \), \( \mathbf{e}_2 = (\frac{1}{2}, 0, \frac{1}{2}) \), and \( \mathbf{e}_3 = (\frac{1}{2}, \frac{1}{2}, 0) \), see [19, 28]. As a result, the lattice sites of the \( N \)-site pyrochlore lattice are given by \( \mathbf{R}_{m, \alpha} = \mathbf{R}_m + \mathbf{r}_\alpha \), where \( m = 1, \ldots, N \), \( \alpha = 1, 2, 3, 4 \).

We consider the spin-\( \frac{1}{2} \) isotropic Heisenberg model. Its Hamiltonian has the following form:

\[
H = J \sum_{\langle m, n \rangle} \mathbf{s}_{m, \alpha} \cdot \mathbf{s}_{n, \beta}
\]

\[
= J \sum_m \left( \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,3} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,4} + \mathbf{s}_{m,2} \cdot \mathbf{s}_{m,3} + \mathbf{s}_{m,2} \cdot \mathbf{s}_{m,4} + \mathbf{s}_{m,3} \cdot \mathbf{s}_{m,4} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2,1,2,1} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2,1,2,3} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2,2,3,3} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2,2,3,1} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2,2,1,3} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2,2,1,1} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2,2,1,2} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2,2,1,3} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2,2,1,1} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2,2,1,2} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2,2,1,3} + \mathbf{s}_{m,1} \cdot \mathbf{s}_{m,2,2,1,1} \right),
\]  

(2.1)

where the first sum in equation (2.1) runs over all nearest neighbors \( m; \alpha \) and \( n; \beta \), \( J = -|J| < 0 \) is the ferromagnetic exchange interaction constant, and the sum in the second line in equation (2.1) runs over all \( m = (m_1, m_2, m_3) \), \( m_\alpha = 1, \ldots, \mathcal{L}_\alpha \) (\( \alpha = 1, 2, 3 \)), \( \mathcal{L}_1 \mathcal{L}_2 \mathcal{L}_3 = N \). The spin operators at different sites commute and satisfy the following commutation relations at the same site:

\[
[s^x, s^z] = 2s^y, \quad [s^y, s^z] = 0, \quad [s^z, s^x] = 2s^y,
\]

(2.2)

where \( s^\pm = s^x \pm is^y \). We have assumed \( \hbar = 1 \) for brevity.
We begin with a brief description of the linear spin-wave theory which should be valid in the low-temperature limit. Using the Holstein-Primakoff transformation
\[ s^+ = \sqrt{2s} \sqrt{1 - \frac{a^+ a}{2s}} a \approx \sqrt{2s} a, \quad s^- = \sqrt{2s} a^\dagger \sqrt{1 - \frac{a^+ a}{2s}} = \sqrt{2s} a^+, \quad s^z = s - \frac{1}{2} a^+ a, \]  
(2.3)
we introduce the bosonic operators \( a_{\mathbf{m}, \alpha} \) (and \( a^\dagger_{\mathbf{m}, \alpha} \)) with \( \alpha = 1, 2, 3, \ldots \) and \( \mathbf{m} = 1, \ldots, N \). Furthermore, we introduce the bosonic operators
\[ a_{\mathbf{q}, \alpha} = \frac{1}{\sqrt{N}} \sum_{\mathbf{m}} e^{-i\mathbf{q} \cdot \mathbf{R}_m} a_{\mathbf{m}, \alpha}, \]  
(2.4)
where \( \mathbf{q} \cdot \mathbf{R}_m = q_1 m_1 + q_2 m_2 + q_3 m_3 \) and \( q_\alpha = 2\pi z_\alpha / L_\alpha, \quad z_\alpha = 1, \ldots, L_\alpha \) (\( \alpha = 1, 2, 3 \)). [Note that \( q_1 = \frac{1}{2}(q_y + q_z), q_2 = \frac{1}{2}(q_x + q_z), \) and \( q_3 = \frac{1}{2}(q_x + q_y) \).] Then, the spin Hamiltonian given in equation (2.1) can be approximately rewritten in the following form:
\[ H = 12N s^2 J - 2s J \sum_{\mathbf{q}} \left( a_{\mathbf{q}, 1}^\dagger a_{\mathbf{q}, 1} a_{\mathbf{q}, 2}^\dagger a_{\mathbf{q}, 2} a_{\mathbf{q}, 3}^\dagger a_{\mathbf{q}, 3} a_{\mathbf{q}, 4}^\dagger a_{\mathbf{q}, 4} \right) \mathbf{F} \]  
(2.5)
where
\[ \mathbf{F} = \begin{pmatrix} 3 & -\phi_1 & -\phi_2 & -\phi_3 \\ -\phi_1^* & 3 & -\phi_{12}^* & -\phi_{13}^* \\ -\phi_2^* & -\phi_{12}^* & 3 & -\phi_{23}^* \\ -\phi_3^* & -\phi_{13}^* & -\phi_{23}^* & 3 \end{pmatrix}, \quad \phi_\alpha = e^{-i\mathbf{q} \cdot \mathbf{a}_\alpha} \cos \frac{\alpha \pi}{2}, \quad \phi_{\alpha \beta} = e^{i\frac{(\alpha - \beta) \mathbf{a}_\alpha \cdot \mathbf{a}_\beta}{2}} \cos \frac{\alpha \pi - \beta \pi}{2}. \]  
(2.6)
After diagonalizing the bilinear Bose form in equation (2.5) we arrive at
\[ H = 12N s^2 J - 2s J \sum_{\gamma=1}^{4} \sum_{\mathbf{q}} \omega_{\gamma, \mathbf{q}} \xi_{\gamma, \mathbf{q}}^x \xi_{\gamma, \mathbf{q}}^y. \]  
(2.7)
The energies of the linear spin waves (magnons) \(-2sJ\omega_{\gamma, \mathbf{q}} = |J|\omega_{\gamma, \mathbf{q}} \) are determined by the eigenvalues of the matrix \( \mathbf{F} \)
\[ \omega_{1, \mathbf{q}} = \omega_{2, \mathbf{q}} = 4, \quad \omega_{3, \mathbf{q}} = 2 + D_\mathbf{q}, \quad \omega_{4, \mathbf{q}} = 2 - D_\mathbf{q}, \]  
(2.8)
The calculated spin-wave dispersion coincides with the result reported in [26] [see figure 1 (c) of that paper].

Now, from equation (2.7) we can obtain the internal (intrinsic) energy per site
\[ \frac{\langle H \rangle}{N} = 3s^2 J - \frac{s J}{2N} \sum_{\gamma=1}^{4} \sum_{\mathbf{q}} \omega_{\gamma, \mathbf{q}} n_{\gamma, \mathbf{q}}. \]  
(2.9)
and the magnetization per site
\[ \langle s^z \rangle = \frac{1}{N} \sum_{\mathbf{m}, \alpha} \langle s_{\mathbf{m}, \alpha}^z \rangle = s - \frac{1}{4N} \sum_{\gamma=1}^{4} \sum_{\mathbf{q}} n_{\gamma, \mathbf{q}}. \]  
(2.10)
Here, \( n_{\gamma, \mathbf{q}} = 1/(e^{\omega_{\gamma, \mathbf{q}}/k_B T} - 1) \) denotes the Bose-Einstein distribution function (we set \( k_B = 1 \) for brevity) and \( s = \frac{1}{2} \). Below we use the linear spin-wave theory predictions (2.9) and (2.10) for comparison with the Green’s function approach results.
3. Green’s function method. Random-phase approximation

Now, we turn to the Green’s function approach [3–5, 11, 22, 32]. We introduce the following operators

\[ s_{q,\alpha}^+ = \frac{1}{\sqrt{\mathcal{V}}} \sum_m e^{iq \cdot R_m} s_{m,\alpha}^+ \]  

(3.1)

and the (retarded) Green’s function

\[ G_{\alpha\beta}(t) \equiv \langle \langle s_{q,\alpha}^+ | s_{q,\beta}^- \rangle \rangle_t \equiv -i \Theta(t) \langle [s_{q,\alpha}^+(t), s_{q,\beta}^-] \rangle, \quad G_{\alpha\beta}(\omega) = \int_{-\infty}^{\infty} dt e^{i\omega t} G_{\alpha\beta}(t), \]  

(3.2)

see, e.g., [29]. The first-order equation of motion reads:

\[ \omega \langle \langle s_{q,\alpha}^+ | s_{q,\beta}^- \rangle \rangle_\omega = \frac{2}{\mathcal{V}} \sum_m \langle s_{m,\alpha}^- \rangle \delta_{\alpha\beta} + i\langle [s_{q,\alpha}^+ | s_{q,\beta}^-] \rangle_\omega, \quad \dot{s}_{q,\alpha}^+ = \frac{1}{\sqrt{\mathcal{V}}} \sum_m e^{-iq \cdot R_m} [s_{m,\alpha}, H]. \]  

(3.3)

Calculating the commutator in the right-hand side of the second equation in equation (3.3) (see appendix) and introducing the random-phase (or Tyablikov) approximation \[ s_{q,\alpha}^+ s_{q}^+ \rightarrow \langle s^2 \rangle s_{q}^+ \] (see appendix) we obtain the closed-form equation for the Green’s function

\[ \sum_{\gamma} (\omega \delta_{\alpha\gamma} + 2J \langle s^2 \rangle F_{\alpha\gamma}) G_{\gamma\beta}(\omega) = 2\langle s^2 \rangle \delta_{\alpha\beta}, \]  

(3.4)

where \(-2J\langle s^2 \rangle F\) is the frequency matrix, \(G(\omega)\) is the matrix of Green’s functions, and \(M = 2\langle s^2 \rangle 1\) is the moment matrix, see also equation (2.8). The matrix \(F\) appeared already in the linear spin-wave calculations, see equation (2.6).

We can find the eigenvalues and the eigenvectors of the matrix \(F\), which satisfy \(\sum_{\beta} F_{\alpha\beta}(\beta \mid \gamma q) = \omega_\gamma q(\alpha \mid \gamma q)\). The eigenvalues \(\omega_\gamma q, \gamma = 1, 2, 3, 4\) are given in equation (2.8). The (orthonormal) eigenvectors \(\langle \alpha \mid \gamma q \rangle\), \(\gamma = 1, 2, 3, 4\) are too lengthy to be presented here.

Finally, using the identity \(1 = \sum_{\gamma} \langle \gamma q \rangle \langle \gamma q \rangle\), we can resolve equation (3.4) with respect to \(G\). \(G = 2\langle s^2 \rangle(\omega 1 + 2J \langle s^2 \rangle F)^{-1} \sum_{\gamma} \langle \gamma q \rangle \langle \gamma q \rangle = \sum_{\gamma}[2\langle s^2 \rangle/(\omega + 2J \langle s^2 \rangle \omega_\gamma q)]\langle \gamma q \rangle \langle \gamma q \rangle\) to get the desired result

\[ G_{\alpha\beta}(\omega) = 2\langle s^2 \rangle \sum_{\gamma=1}^{4} \frac{\langle \alpha \mid \gamma q \rangle \langle \gamma q \mid \beta \rangle}{\omega + 2J \langle s^2 \rangle \omega_\gamma q}. \]  

(3.5)

Here, \(\langle \gamma q \mid \beta \rangle\) is the \(\beta\)-th component of the eigenvector \(\langle \gamma q \rangle\). Clearly, the magnetization \(\langle s^2 \rangle\) which enters equation (3.5) should be determined self-consistently, see equation (3.7) below.

As can be seen from equation (3.5), the excitation energies are given by \(-2J\langle s^2 \rangle \omega_\gamma q = 2J[\langle s^2 \rangle \omega_\gamma q].\) In the limit \(T \rightarrow 0\), when \(\langle s^2 \rangle \rightarrow \frac{1}{2}\), they coincide with the magnon energies considered in section 2, and therefore, we call these excitations magnons. Moreover, they coincide with the results of [19] in this limit (see the upper panel in figure 2 of that paper). For finite temperatures \(T > 0\) the dependences on \(q\) remain unchanged, and the energies simply decrease due to the factor \(\langle s^2 \rangle < \frac{1}{2}\). More sophisticated effects of the temperature on the excitation energies were discussed within the rotation-invariant (or Kondo-Yamaji [33]) Green’s function approach in [19]. Since \(D_q \rightarrow 2 - \frac{1}{\rho^2} q^2\) as \(q \rightarrow 0\), \(\omega_{q,q}\) is the acoustic branch of the spectrum, i.e., \(\omega_{q,q} \rightarrow \rho |q|^2, \quad \rho = \frac{1}{16} 4J [\langle s^2 \rangle] q \rightarrow 0\). The spin stiffness \(\rho\) is proportional to \(\langle s^2 \rangle\); it shows the same temperature behavior as \(\langle s^2 \rangle\) vanishing as \(T \rightarrow T_c\).  

The rotation-invariant Green’s function approach considers the equation of motion up to the second order, which obviously involves Green’s functions of higher order than the initial ones. Several products of three-spin operators are then simplified by a decoupling like \(s_{q,\alpha}^+ s_{q}^+ \rightarrow \alpha_{AB} s_{q,\alpha}^+ s_{q}^+\) etc., where \(\alpha_{AB}\) is a vertex parameter which is introduced to improve the approximation made by decoupling. After all, the correlation functions, the vertex parameter, and the condensation term which is related to magnetic long-range order are determined self-consistently within the Tyablikov approximation, one face only one resulting equation for \(\langle s^2 \rangle\), see equation (2.7). For further details see [19].
Knowing the Green’s functions, we calculate the equal-time correlation functions via the relation

\[
\langle \phi_{q,\alpha}^{+} \phi_{q,\alpha} \rangle = \frac{1}{2\pi} \lim_{\varepsilon \rightarrow 0} \int_{-\infty}^{\infty} d\omega \frac{G_{\alpha\beta}(\omega + i\varepsilon) - G_{\alpha\beta}(\omega - i\varepsilon)}{e^{\frac{\omega}{T}} - 1} = 2 \langle s^{z} \rangle \sum_{\gamma = 1}^{4} \frac{\langle \alpha | q | \gamma \rangle \langle q | \gamma | \beta \rangle}{e^{-\frac{2J(q^2)/\omega_{i\gamma,q}}{\omega - 1}}}. \tag{3.6}
\]

These correlation functions yield important thermodynamic characteristics. In particular, using the identity \(s^{z} = \frac{1}{2} - s^{-} s^{+} \) which leads to \(\langle s^{z} \rangle = \frac{1}{2} - \frac{1}{N} \sum_{m,n} \langle s_{m,n}^{z} s_{m,n}^{+} \rangle = \frac{1}{2} - \frac{1}{N} \sum_{n} \sum_{q} \langle s_{n,q}^{z} s_{q,n}^{+} \rangle \), we arrive at the following formula for \(\langle s^{z} \rangle\):

\[
\langle s^{z} \rangle = \frac{1}{2} - \frac{1}{2N} \sum_{q = 1}^{4} \frac{1}{\sum_{\gamma = 1}^{4} e^{-\frac{2J(q^2)/\omega_{i\gamma,q}}{\omega - 1}}}. \tag{3.7}
\]

In fact, this is the equation for determining self-consistently the magnetization \(\langle s^{z} \rangle\). We can obtain the internal energy of the spin model at hand by averaging the Hamiltonian \((H)\) over the magnetic phase above the critical (Curie) temperature \(T_{c}\).

Since we have calculated only the Green’s function \(\langle \langle s^{z} | s^{z} \rangle \rangle\) rather than \(\langle \langle s^{z} | s^{z} \rangle \rangle\) we should eliminate the correlations \(\langle s^{z} s^{z} \rangle\) from the consideration in order to treat all correlations in \((H)\) on equal footing, i.e., within the same (Tyablikov) approximation \([31, 34]\). After some manipulations (see appendix), we arrive at the following expression for the internal energy within the Tyablikov approximation:

\[
\frac{\langle H \rangle}{N} = \frac{3}{2} \langle J \rangle \langle s^{z} \rangle - \frac{\langle J \rangle \langle s^{z} \rangle^{2}}{2N} + \sum_{y = 1}^{4} \sum_{q} \frac{\omega_{y,q}}{e^{\frac{\omega_{y,q}}{T}} - 1} + \frac{\langle J \rangle \langle s^{z} \rangle}{2N} \sum_{y = 1}^{4} \sum_{q} \sum_{r} \phi_{y,r} \langle \alpha | q | \gamma \rangle \langle q | \gamma | \beta \rangle e^{-\frac{2J(q^2)/\omega_{i\gamma,q}}{\omega - 1}},
\]

where \(\phi_{y,1}^{1} = \varepsilon_{1}, \phi_{y,1}^{2} = \varepsilon_{13}, \phi_{y,2} = \phi_{y,1}, \phi_{y,1}^{3} = \phi_{y,1}^{4} = \phi_{y,1}, \phi_{y,1}^{4} = \phi_{y,1}^{3}, \phi_{y,1}^{3} = \phi_{y,1}^{2}, \phi_{y,1}^{2} = \phi_{y,1}^{1} \).

Differentiation of the internal energy with respect to \(T\) gives the specific heat.

The Green’s function \((3.5)\) gives the dynamic susceptibility \(\chi_{q}^{+}(\omega) \equiv \frac{1}{4} \sum_{\alpha,\beta = 1}^{N} \chi_{q,\alpha\beta}^{+}(\omega)\), \(\chi_{q,\alpha\beta}^{+}(\omega) = -G_{\alpha\beta}(\omega)\), which is related to the dynamic structure factor

\[
\delta_{q}^{\pm}(\omega) = \frac{2}{1 - e^{-\frac{\omega}{T}}} \text{Im} \chi_{q}^{\pm}(\omega) = \frac{\pi \langle s^{z} \rangle}{1 - e^{-\frac{\omega}{T}}} \sum_{\alpha,\beta = 1}^{4} \langle \alpha | q | \gamma \rangle \langle q | \gamma | \beta \rangle \delta (\omega - 2J \langle s^{z} \rangle \omega_{i\gamma,q}). \tag{3.9}
\]

Furthermore, the Green’s function given in equation \((3.5)\) is proportional to \(\langle s^{z} \rangle\) and at zero field it disappears in the high-temperature paramagnetic phase above the critical (Curie) temperature \(T_{c}\). However, the (initial) susceptibility \(\chi\), defined as \(\langle s^{z} \rangle = \chi h\), where \(h\) is an infinitesimally small applied magnetic field can be determined as follows. For the spin model with the Hamiltonian \(H = -h \sum_{m,n} s_{m,n}^{z}\), the Green’s function \(G_{\alpha\beta}(\omega)\) within the random-phase approximation is again given by equation \((3.5)\), although with \(2J \langle s^{z} \rangle \omega_{i\gamma,q} - h\) instead of \(2J \langle s^{z} \rangle \omega_{i\gamma,q}\) in the denominator. Therefore, the equation for \(\langle s^{z} \rangle\) now reads

\[
\langle s^{z} \rangle = \frac{1}{2} - \frac{\langle s^{z} \rangle}{2N} \sum_{y = 1}^{4} \sum_{q} \frac{1}{\sum_{\gamma = 1}^{4} e^{-\frac{2J(q^2)/\omega_{i\gamma,q} - h}{\omega - 1}}}, \tag{3.10}
\]

cf. equation \((3.7)\). Following, e.g., \([29]\), for \(T > T_{c}\) we set \(\langle s^{z} \rangle = 0\) in the left-hand side of equation \((3.10)\) and then expand the exponent and substitute \(\langle s^{z} \rangle = \chi h\) to arrive at the equation for \(1/\chi\) above the critical temperature \(T_{c}\):

\[
1 = \frac{1}{\chi} \sum_{y = 1}^{4} \sum_{q} \frac{T}{2J \omega_{i\gamma,q} + \frac{1}{T}}. \tag{3.11}
\]

All calculations described in this section are, in principle, well known \([3, 5, 11, 29, 32]\). However, to our best knowledge, we are not aware of any random-phase Green’s function paper for the pyrochlore-lattice quantum Heisenberg ferromagnet.
4. Low- and intermediate-temperature results

In the low-temperature limit, the excitations described by $G_{\alpha\beta}(\omega)$ \(3.2, 3.3\) are magnons also emerging in the linear spin-wave theory. Consequently, in the low-temperature limit, the Green’s function results are expected to coincide with the ones of the linear spin-wave theory, see the low-temperature region in figures 1 and 2.

Let us compute the critical (Curie) temperature $T_c$. At the critical temperature, $\langle s^z \rangle$ vanishes; we set $\langle s^z \rangle = 0$ in the left-hand side of equation (3.7) and expand the exponent in the right-hand side of equation (3.7) to obtain the critical temperature:

$$
\frac{T_c}{|J|} = \frac{2}{\sum_{\alpha=1}^{4} \sum_{\omega_{\mu\nu}} 1} \approx 0.872.
$$

(4.1)

Here, in the thermodynamic limit $\frac{1}{\chi} \sum_{q} \ldots \rightarrow \frac{1}{\pi} \int_{0}^{\pi} dq_{1} \int_{0}^{\pi} dq_{2} \int_{0}^{\pi} dq_{3} \ldots$. The same equation follows from equation (3.11) after setting $1/\chi = 0$. This result should be compared to the corresponding ones obtained by other means given in the second column of table 1. The important feature is visible: the critical temperature calculated within the random-phase approximation Green’s function method is the highest one compared with the results of other approaches (excluding mean-field) indicating some underestimate of the role of thermal fluctuations. Comparing the results for the pyrochlore and simple-cubic cases in the same approximation, we again observe a manifestation of finite-temperature frustration effects \(19, 35\): thermal fluctuations destroy the magnetic order more effectively for the pyrochlore-lattice geometry, and the critical temperature for the pyrochlore ferromagnet is about 15% smaller than for the simple-cubic ferromagnet. Of course, the mean-field result for both lattices is the same: $T_c = \frac{2}{J}$.

We can solve equation (3.7) with respect to $\langle s^z \rangle$ for all temperatures below $T_c$ (4.1). This temperature dependence of the magnetization is shown in figure 1 along with the results of the rotation-invariant Green’s function method \(19\) and the quantum Monte Carlo simulations using the ALPS package \(34, 37\) (see also \(38\)). We also report in figure 1 the corresponding results for the simple-cubic case for comparison. Moreover, in figure 1 we show additionally the temperature dependence of $1/\chi$ in the paramagnetic phase as it follows from equation (3.11) (thick solid), from the rotation-invariant

![Figure 1](image-url)

*Figure 1.* (Colour online) Temperature dependence of magnetization $m = \langle s^z \rangle$ (thin) and inverse susceptibility $1/\chi$ (thick) for the spin-$\frac{1}{2}$ pyrochlore-lattice (red) Heisenberg ferromagnet ($J = -1$) obtained within Tjablikov approximation (solid) and rotation-invariant Green’s function method (dashed). For comparison, we also show the results for the simple-cubic lattice (black). Note that the mean-field data (solid green) are identical for both lattices. Quantum Monte Carlo simulation data for $N = 16 \cdot 32^3$ sites (pyrochlore, red) and $N = 80^3$ sites (simple-cubic, black) are shown by open circles ($m$) and filled circles ($1/\chi$). Inset: Magnetization at low temperatures. We also show here the linear spin-wave theory predictions (very thin dotted).
The spin-1/2 Heisenberg ferromagnet on the pyrochlore lattice

We compare the results for the pyrochlore lattice (red) and the simple-cubic lattice (black) which are identical at the mean-field level (solid green). We provide in addition quantum Monte Carlo data obtained for \( N = 16 \cdot 16^3 \) (pyrochlore, red empty circles) and \( N = 40^3 \) (simple-cubic, black empty circles) sites. Very thin dotted (double-dashed) lines represent the 13th order high-temperature expansion raw (extrapolated by the \([6, 7]\) Padé approximant) results.

Inset: Specific heat at low temperatures. We also show here the linear spin-wave theory predictions (very thin dotted). It is in order to mention here, that for the \( c(T) \) curve of the simple-cubic lattice presented in figure 8 of \([19]\), an incorrect data set was used.

|                  | pyrochlore lattice | simple-cubic lattice | \( \frac{T_c^{\text{Pyro}}}{T_c^{\text{Sc}}} \) |
|------------------|--------------------|----------------------|---------------------------------------------|
| QMC              | 0.718 [19]         | 0.839(1) [39]        | 86%                                         |
| HTE              | 0.724…0.754 [40]  | 0.827 [40]           | 88%                                         |
| PFFRG            | 0.77(4) [20]       | 0.90(4) [20]         | 86%                                         |
| RGM              | 0.778 [19]         | 0.926 [38, 41]       | 84%                                         |
| RPA              | 0.872 (this paper) | 0.989 [29]           | 88%                                         |
| MFA              | \( \frac{1}{3} \)  | \( \frac{1}{3} \)    | 100%                                        |

Green’s function method \([19]\) (thick dashed), and from the quantum Monte Carlo simulations (filled circles) for the pyrochlore (red) and simple-cubic (black) lattices. Within the mean-field approximation \( 1/\chi = 4(T_c - T) \), \( T_c = \frac{3}{2}|J| \) for both lattices, see thick solid green curve. The reported results for the temperature dependences of magnetization and susceptibility agree well with each other and illustrate the finite-temperature frustration effects as well as the quality of various approximations.

The specific heat within the Tyablikov approximation is evaluated by finding the derivative of equation (3.8) with respect to temperature \( T \). In figure 2 we compare the temperature dependence of the specific heat obtained by different methods for the pyrochlore lattice (red) and simple-cubic lattice.
pyrochlore-lattice Heisenberg ferromagnet ($J = -1$) as a function of the reduced momentum $t = 2 - D_q$ with $q = (q, q, q)$ at $T = 0.0425$ obtained within the Tyablikov approximation. $\delta$-function is smeared by introducing the Lorentzian function with finite $\epsilon = 0.01$.

The mean-field result for both lattices is the same (green): the critical temperature $T_c$ is overestimated and $c(T_c)$ is finite. The Tyablikov approximation (solid red and solid black) yields already different values for $T_c$, although the specific heat diverges at the Curie temperature. This contradicts the current critical exponent estimates of the three-dimensional Heisenberg model obtained by field theory [42], high-temperature series analyses, and Monte Carlo [43–45] methods which predict that the specific heat is not singular for this model. According to the rotation-invariant Green’s function method, $c(T_c)$ is finite (dashed red and dashed black). We also report quantum Monte Carlo data obtained using the ALPS package [36, 37] (see also [38]); red (black) empty circles refer to the pyrochlore (simple-cubic) lattice of $N = 16 \cdot 16^3 (N = 40^3)$ sites. Moreover, we show the 13th order high-temperature expansion results, raw data (very thin dotted) and [22] Padé approximants (very thin double-dashed), obtained using the HTE program freely available at http://www.uni-magdeburg.de/jschulen/HTE, see [40, 46]. In summary, the presented data show that the Tyablikov approximation is capable of reproducing reasonably well the dependence $c(T)$ in the ferromagnetic phase, except the region close to $T_c$. In the paramagnetic phase, the rotation-invariant Green’s function method and, of course, high-temperature expansion method give reasonably good results.

Finally, we turn to the dynamic structure factor $S_{q}^{+\to-}(\omega)$. Within the Tyablikov approximation, it is given by equation (3.9). For further calculations we replace in equation (3.9) $\delta(x)$ by the Lorentzian function $\epsilon/[\pi(x^2 + \epsilon^2)]$ with $\epsilon = 0.01$. Inspired by the experimental paper on the spin-$\frac{1}{2}$ pyrochlore ferromagnet $\text{Lu}_2\text{V}_2\text{O}_7$ [26], where neutron inelastic scattering data were reported (see figure 2 of that paper), we calculate $S_{q}^{+\to-}(\omega)$ at $T = 0.0425|J|$ along the path $q = (q, q, q)$ and present the result as a function of the reduced momentum $t = 2 - D_q$, see figure 3. (Experimentalists collected data from more points in the $q$-space, not only along the path $q = (q, q, q)$, see [26].) Overall, figure 3 resembles the experimental data reported in [26]: the dynamic structure factor is concentrated along the lines $\omega_{kq} = 4 - t$, $\omega_{3q} = t$, however, because of the chosen path $q = (q, q, q)$, not along the line $\omega_{1q} = \omega_{2q} = 4$. While comparing with experiments (and with the rotation-invariant Green’s function result [19]), one should remember that $S_{q}^{+\to-}(\omega)$ presents only the transverse part, while the longitudinal part $S_{q}^{zz}(\omega)$ is not taken into consideration. We note in passing that comparing theoretical predictions with experimental data one can obtain the exchange interaction constants, and the exploited here random-phase approximation Green’s function method can be used for this purpose.

5. Summary

In the present paper we have discussed the thermodynamic properties of the pyrochlore-lattice quantum Heisenberg ferromagnet. Our main results (Curie temperature, temperature dependences of the magnetization and the specific heat, dynamic structure factor) have been obtained within the random-
phase approximation. We compare them to the linear spin-wave theory at low temperatures as well as to the rotation-invariant Green’s function method \[19\], quantum Monte Carlo simulation, and high-temperature expansion \[12\, 20\] results. Although the random-phase approximation goes beyond the mean-field treatment, thus leading to different results for the pyrochlore-lattice and the simple-cubic-lattice case, it takes into account not all fluctuations, and ferromagnetic ordering is slightly overestimated in comparison with the second-order rotation-invariant Green’s function method results. Nevertheless, the random-phase approximation Green’s function method together with the high-temperature expansion provide a simple route to describe quantum Heisenberg ferro magnets for all temperatures. Of course, the quantum Monte Carlo simulations or the rotation-invariant Green’s function method are also applicable, though these are more complicated techniques.

The Green’s function method which uses the Tyablikov approximation can be applied to some other spin models related to the pyrochlore lattice. For example, one can in a similar way consider the ferromagnet in the presence of the breathing anisotropy or the second-nearest neighbor interactions. However, the most interesting problem within this context is the study of the properties of the pyrochlore-lattice ferro- and antiferromagnet are connected by the obvious relation: \( Z_{\text{FM}}(-T) = Z_{\text{AFM}}(T) \). This implies that the specific heat of the antiferromagnet at temperature \( T \) is given by the specific heat of the ferromagnet taken at temperature \(-T\). However, in the thermodynamic limit, because of spontaneous symmetry breaking, this relation is not valid. Clearly, the rotation-invariant Green’s function approach, which implies \( \langle s^z \rangle = 0 \), seems to be an appropriate approximation for examining that case. The work in this direction is in progress \[47\].

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**A. First-order equation of motion, Tyablikov approximation, and some other calculations**

To obtain the first-order equation of motion we have to calculate the commutator in the second equation in equation (3.3). We begin with the case \( \alpha = 1 \) in equation (3.3). Then,

\[
\frac{1}{J} \left[ \mathbf{s}_{m1}^+, \mathbf{H} \right] = \sum_n \left[ \frac{1}{2} s_{n1}^+ \left( s_{n2}^+ + s_{n3}^+ + s_{n4}^+ \right) + s_{n1}^z \left( s_{n2}^z + s_{n3}^z + s_{n4}^z \right) \right. \\
\left. + \frac{1}{2} s_{n1}^z \left( s_{n1-n2,n3}^z + s_{n1-n2,n3}^z + s_{n1,n2-n3}^z \right) + s_{n1}^z \left( s_{n1-n2,n3}^z + s_{n1,n2-n3}^z + s_{n1,n2-n3}^z \right) \right]
\]

\[
\begin{align*}
&= s_{m1}^z \left( s_{m2}^+ + s_{m3}^+ + s_{m4}^+ \right) - s_{m1}^z \left( s_{m2}^z + s_{m3}^z + s_{m4}^z \right) \\
&\quad + s_{m1}^z \left( s_{m1-m2,m3}^z + s_{m1,m2-1,m3}^z + s_{m1,m2,m3}^z \right) \\
&\quad - s_{m1}^z \left( s_{m1-m2,m3}^z + s_{m1,m2-1,m3}^z + s_{m1,m2,m3}^z \right) .
\end{align*}
\]  

(A.1)

Within the Tyablikov approximation \( s^z s^z \rightarrow \langle s^z \rangle s^z \) (obviously, two spin operators here are attached to different sites and \( \langle s^z \rangle \) is already site independent) and, therefore,

\[
\frac{1}{\sqrt{N}} \sum_m e^{-i q \mathbf{R}_m} \left[ s_{m1}^+, \mathbf{H} \right] \rightarrow -2J \langle s^z \rangle \left( 3s_{q,1}^z - \phi_1 s_{q2}^z - \phi_2 s_{q3}^z - \phi_3 s_{q4}^z \right) ;
\]  

(A.2)

for the definition of \( \phi \), see equation (2.6). As a result, according to equation (3.3), we obtain...
\[ \omega G_{1p}(\omega) = 2(s^2)\delta_{1p} - 2J(s^2) \left[ 3G_{1p}(\omega) - \phi_1 G_{2p}(\omega) - \phi_2 G_{3p}(\omega) - \phi_3 G_{4p}(\omega) \right]. \] (A.3)

Setting \( \alpha = 2, \alpha = 3, \) and \( \alpha = 4 \) and repeating such calculations thrice, we obtain three more equations. After all, we may combine them in the matrix form

\[ (\omega I + 2J(s^2)F)\Omega(\omega) = 2(s^2)\Omega, \] (A.4)

where the matrix \( F \) is defined in equation (3.6), cf. equation (3.4).

It is worth commenting briefly on the mean-field approximation. Within the frames of this approximation we replace equation (2.1) by \( H = -3NJ(s^2)^2 + 6J(s^2)\sum m(s^2_m + s^2_m + s^2_m + s^2_m). \) After that, the exact first-order equation (3.3) \( \omega G_{\alpha p}(\omega) = 2(s^2)\delta_{\alpha p} - 6J(s^2)G_{\alpha p}(\omega) \) immediately yields \( G_{\alpha p}(\omega) = 2(s^2)\delta_{\alpha p}/(\omega + 6J(s^2)) \) that results in \( \langle \omega s^2\rangle \rangle = 2(s^2)\delta_{\alpha p}/(\omega + 6J(s^2)/T - 1) \), cf. equations (3.5) and (3.6). The self-consistent equation for \( \langle s^2 \rangle \) reads: \( 1 = 2(s^2)\coth(3J\langle s^2 \rangle /T) \) and it yields \( T = \beta J \), cf. equations (3.7) and (4.1). For the internal energy, we have: \( \langle H \rangle = -3J\langle s^2 \rangle^2 \), cf. equation (3.8). The inverse susceptibility above \( T_c \) is as follows: \( 1/J = \frac{\ln(1)}{4(T_c - T)} \), cf. equation (3.11). Clearly, the mean-field results for observables are identical for the pyrochlore lattice and for the simple-cubic lattice.

Let us explain how to calculate the internal energy within the Tyablikov approximation. The internal energy is defined as the average of Hamiltonian (2.1), i.e., as \( \langle H \rangle \). To account the transverse correlation energy and the longitudinal correlation energy on equal footing, let us obtain first a formula for \( \langle H \rangle \) that will contain only such correlation functions which are evaluated straightforwardly using the same Tyablikov approximation. We begin with calculating the commutators \( [s^+_{n\alpha}, H], \alpha = 1, 2, 3, 4, \) see equation (A.1), which yield the correlation functions \( \langle s^+_{n\alpha}[s^+_{n\alpha}, H] \rangle, \alpha = 1, 2, 3, 4. \) We use the relations \( s^- s^+ = s^- s^+ s^+ + s^- s^+ = \frac{1}{2} - s^2 \) to arrive at the wanted identity with \( \langle H \rangle \):

\[
\begin{align*}
\frac{4}{N} \sum_{\alpha=1}^{N} \langle s^+_{n\alpha}[s^+_{n\alpha}, H] \rangle = & -12NJ\langle s^2 \rangle + 2\langle H \rangle \\
+ & J \sum_{n} \left[ \langle (s^- s^+_{q,1}, s^+_{q,3}) \rangle (1 + e^{-i\eta}) + \langle (s^- s^+_{q,2}, s^+_{q,3}) \rangle (1 + e^{-i\eta}) + \langle (s^- s^+_{q,1}, s^+_{q,4}) \rangle (1 + e^{-i\eta}) \\
+ & \langle (s^- s^+_{q,2}, s^+_{q,4}) \rangle (1 + e^{-i\eta}) + \langle (s^- s^+_{q,1}, s^+_{q,2}) \rangle (1 + e^{i\eta}) + \langle (s^- s^+_{q,3}, s^+_{q,2}) \rangle (1 + e^{i\eta}) \\
+ & \langle (s^- s^+_{q,3}, s^+_{q,4}) \rangle (1 + e^{i\eta}) + \langle (s^- s^+_{q,4}, s^+_{q,2}) \rangle (1 + e^{i\eta}) + \langle (s^- s^+_{q,4}, s^+_{q,3}) \rangle (1 + e^{i\eta}) \right] \\
= & -12NJ\langle s^2 \rangle + 2\langle H \rangle + 2J \sum_{q} \left[ \langle (s^- s^+_{q,1}, s^+_{q,2}) \rangle \phi_{1} + \langle (s^- s^+_{q,1}, s^+_{q,3}) \rangle \phi_{2} + \langle (s^- s^+_{q,1}, s^+_{q,4}) \rangle \phi_{3} \\
+ & \langle (s^- s^+_{q,2}, s^+_{q,1}) \rangle \phi_{4} + \langle (s^- s^+_{q,2}, s^+_{q,3}) \rangle \phi_{5} + \langle (s^- s^+_{q,2}, s^+_{q,4}) \rangle \phi_{6} + \langle (s^- s^+_{q,3}, s^+_{q,2}) \rangle \phi_{7} \\
+ & \langle (s^- s^+_{q,3}, s^+_{q,4}) \rangle \phi_{8} + \langle (s^- s^+_{q,4}, s^+_{q,2}) \rangle \phi_{9} + \langle (s^- s^+_{q,4}, s^+_{q,3}) \rangle \phi_{10} \right];
\end{align*}
\] (A.5)

(To keep notations short, we have suppressed some site indexes) or using the \( q \)-momentum representation, see equation (3.11).
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for definition of \(\phi_a\) and \(\phi_{ab}\) see equation (3.6). In the left-hand side of equation (A.6) we can obviously write

\[
\langle s_{\alpha}^+ [s_{\alpha}^-, H]\rangle = i \left[ \frac{d}{dt} \langle s_{\alpha}^- s_{\alpha}^+(t) \rangle \right]_{t=0} = i \left[ \frac{d}{dt} \frac{1}{2\pi} \int_{-\infty}^{\infty} d\omega e^{-i\omega t} J_{\alpha\alpha}(\omega) \right]_{t=0} = \frac{1}{2\pi} \int_{-\infty}^{\infty} d\omega \omega J_{\alpha\alpha}(\omega),
\]

(A.7)

where

\[
J_{\alpha\alpha}(\omega) = i G_{\alpha\alpha}(\omega + i0) - G_{\alpha\alpha}(\omega - i0)
\]

\[
e^{-\frac{\omega}{\Theta}} - 1
\]

(A.8)

we have used equation (3.5). That is,

\[
\sum_{\alpha} \sum_{\mathbf{q}} \langle s_{\alpha}^- [s_{\alpha}^+, H]\rangle = 4|J| \langle s^z \rangle^2 \sum_{\gamma=1}^{4} \sum_{\mathbf{q}} \frac{\omega_{\gamma\mathbf{q}}}{e^{\frac{\omega_{\gamma\mathbf{q}}}{\Theta}} - 1}.
\]

(A.9)

In the right-hand side of equation (A.6) we have the correlation functions \(\langle (s^z s^z)_{\gamma\beta} s_{\alpha\alpha}^+ \rangle\) which are calculated from the Green’s function \(H_{\alpha\beta}(t) \equiv \langle (s_{\alpha}^+ s_{\alpha}^-)_{\gamma\beta} \rangle \rangle = -i\Theta(t)\langle (s_{\alpha}^+ (t), (s^z s^z)_{\gamma\beta}) \rangle\), \(H_{\alpha\beta}(\omega) = \int_{-\infty}^{\infty} dt e^{i\omega t} H_{\alpha\beta}(t)\) using the spectral representation

\[
\langle (s^z s^z)_{\gamma\beta} \rangle = \frac{i}{2\pi} \int_{-\infty}^{\infty} d\omega \frac{H_{\alpha\beta}(\omega + i0) - H_{\alpha\beta}(\omega - i0)}{e^{\frac{\omega}{\Theta}} - 1},
\]

(A.10)

cf. equation (3.6). Repeating the derivation of equation (A.4) (equation of motion within the Tyablikov approximation), but now for the Green’s function \(H(\omega)\), we arrive at

\[
\sum_{\gamma} (\omega_{\gamma\gamma} + 2J \langle s^z \rangle F_{\gamma\gamma}) H_{\gamma\gamma}(\omega) = -\langle s^z \rangle \delta_{\alpha\beta},
\]

(A.11)

cf. equations (3.4) and (A.4), and as a result we obtain

\[
H_{\alpha\beta}(\omega) = -\langle s^z \rangle \sum_{\gamma=1}^{4} \frac{\langle \alpha | q \gamma \rangle \langle q \gamma | \beta \rangle}{\omega + 2J \langle s^z \rangle \omega_{\gamma\mathbf{q}}},
\]

(A.12)

cf. equation (3.6). Therefore,

\[
\langle (s^z s^z)_{\gamma\beta} s_{\alpha\alpha}^+ \rangle = -\langle s^z \rangle \sum_{\gamma=1}^{4} \frac{\langle \alpha | q \gamma \rangle \langle q \gamma | \beta \rangle}{e^{\frac{\omega_{\gamma\mathbf{q}}}{\Theta}} - 1},
\]

(A.13)

cf. equation (3.6). Inserting equation (A.9) and equation (A.13) into equation (A.6), we obtain

\[
4|J| \langle s^z \rangle^2 \sum_{\gamma=1}^{4} \sum_{\mathbf{q}} \frac{\omega_{\gamma\mathbf{q}}}{e^{\frac{\omega_{\gamma\mathbf{q}}}{\Theta}} - 1} = 3N|J| \langle s^z \rangle + 2\langle H \rangle + 2|J| \langle s^z \rangle \sum_{\gamma=1}^{4} \sum_{\mathbf{q}} \frac{\varphi_{\alpha\beta}(\omega_{\gamma\mathbf{q}})}{e^{\frac{\omega_{\gamma\mathbf{q}}}{\Theta}} - 1},
\]

where

\[
\varphi_{12} = \varphi_{21} = \phi_1, \quad \varphi_{13} = \varphi_{31} = \phi_2, \quad \varphi_{14} = \varphi_{41} = \phi_3,
\]

\[
\varphi_{23} = \varphi_{32} = \phi_{12}, \quad \varphi_{24} = \varphi_{42} = \phi_{13}, \quad \varphi_{34} = \varphi_{43} = \phi_{23},
\]

(A.14)

This is equivalent to equation (3.8). Note also that the expression for the internal energy \(\mathcal{U}\) agrees with the formula given in equation (336) of [31].
References

1. Kadanoff L.P., Baym G., Quantum Statistical Mechanics: Green’s Function Methods in Equilibrium and Nonequilibrium Problems, Benjamin, New York, 1962.
2. Abrikosov A.A., Gorkov L.P., Dzyaloshinski I.E., Methods of Quantum Field Theory in Statistical Physics, Prentice-Hall, Englewood Cliffs, N.J., 1963.
3. Zubarev D.N., Sov. Phys. Usp., 1960, 3, 320, doi:10.1070/PU1960v003n03ABEH003275
4. Bonch-Bruevich V.L., Tyablikov S.V., The Green Function Method in Statistical Mechanics, North Holland Publishing Co., Amsterdam, 1962.
5. Elk K., Gasser W., Die Methode der Greenschen Funktionen in der Festkörperphysik, Akademie-Verlag, Berlin, 1979, (in German).
6. Tyablikov S.V., Ukr. Fiz. Zh., 1969, 14, 1097 (in Ukrainian).
7. Stasyuk I.V., Phys. Status Solidi B, 1970, 39, K35, doi:10.1002/pssb.19700390144
8. Stasyuk I.V., Fiz. Met. Metalloved., 1971, 31, 699 (in Russian).
9. Slobodyan P.M., Stasyuk I.V., The Green Function Method in Statistical Mechanics, North Holland Publishing Co., Amsterdam, 1962.
10. Editorial, Condens. Matter Phys., 2008, 11, 401.
11. Stasyuk I.V., Green’s Functions in Quantum Statistics of Solid State: a Textbook, Ivan Franko National University of Lviv, Lviv, 2013, (in Ukrainian).
12. Vladimirov A.A., Ihle D., Plakida N.M., J. Exp. Theor. Phys., 2016, 122, 1060, doi:10.1134/S106377611606008X
13. Mi B.-Z., Solid State Commun., 2016, 239, 20, doi:10.1016/j.ssc.2016.04.015
14. Sun N.-N., Wang H.-Y., J. Magn. Magn. Mater., 2018, 454, 176, doi:10.1016/j.jmmm.2018.01.076
15. Schollwöck U., Richter J., Farnell D.J.J., Bishop R.F. (Eds.), Quantum Magnetism, Lecture Notes in Physics 645, Springer-Verlag, Berlin, 2004.
16. Lacroix C., Mendels P., Mila F. (Eds.), Introduction to Frustrated Magnetism, Springer Series in Solid-State Sciences 164, Springer-Verlag, Berlin, Heidelberg, 2011.
17. Gardner J.S., Gingrich M.J.P., Greedan J.E., Rev. Mod. Phys., 2010, 82, 53, doi:10.1103/RevModPhys.82.53
18. Gao J., Wang H.-Y., Phys. Rev. B, 2017, 96, 174419, doi:10.1103/PhysRevB.96.174419
19. Asai Y., Müller T., Ghosh P., Gingrich M.J.P., Jeschke H.O., Rachel S., Reuther J., Thomale R., Preprint arXiv:1802.09546, 2018.
20. Garcia-Adeva A.J., J. Phys.: Condens. Matter, 2014, 26, 136002, doi:10.1088/0953-8984/26/13/136002
21. Menyuk N., Dwight K., Wold A., J. Appl. Phys., 1966, 37, 1387, doi:10.1063/1.1708484
22. Wojtowicz P.J., IEEE Trans. Magn., 1969, 5, 840, doi:10.1109/TMAG.1969.1066666
23. Yasui Y., Soda M., Ito M., Sato M., Hamaguchi N., Matsushita T., Wada N., Takeuchi T., Aso N., Kakuraki K., J. Phys. Soc. Jpn., 2003, 72, 3014, doi:10.1143/JPSJ.72.3014
24. Yaresko A.N., Phys. Rev. B, 2008, 77, 115106, doi:10.1103/PhysRevB.77.115106
25. Mena M., Perry R.S., Perring T.G., Le M.D., Guerrero S., Sorni M., Adroja D.T., Rüegg Ch., McMorrow D.F., Phys. Rev. Lett., 2014, 113, 047202, doi:10.1103/PhysRevLett.113.047202
26. Tymoshenko Y.V., Onykiienko Y.A., Müller T., Thomale R., Rachel S., Camera A.S., Portnichenko P.Y., Efremov D.V., Tsurkan V., Abernathy D.L., Ollivier J., Schneidewind A., Piovano A., Felea V., Loidl A., Inosov D.S., Phys. Rev. X, 2017, 7, 041049, doi:10.1103/PhysRevX.7.041049
27. URL http://lampx.tugraz.at/~hadley/ss1/bzones/fcc.php
28. Tyablikov S.V., Methods in the Quantum Theory of Magnetism, Plenum Press (Plenum Publishing Corporation), New York, 1967.
29. Gasser W., Heiner E., Greensche Funktionen in Festkörperphysik und Vielteilchenphysik, Wiley-Blackwell, Berlin, 2001, (in German).
30. Fröbrich P., Kuntz P.J., Phys. Rep., 2006, 432, 223, doi:10.1016/j.physrep.2006.07.002
31. Majlis N., The Quantum Theory of Magnetism, World Scientific, Singapore, 2007.
32. Kondo J., Miyashita K., Prog. Theor. Phys., 1972, 47, 807, doi:10.1143/PTP.47.807
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37. Bauer B., Carr L.D., Evertz H.G., Feiguin A., Freire J., Fuchs S., Gamper L., Gukelberger J., Gull E., Guertler S., et al., J. Stat. Mech., 2011, 2011, P05001, doi:10.1088/1742-5468/2011/05/P05001.
38. Menchyshyn O., Krokhmalskii T., Derzhko O., Preprint of the Institute for Condensed Matter Physics, ICMP-14-01E, Lviv, 2014.
39. Wessel S., Phys. Rev. B, 2010, 81, 052405, doi:10.1103/PhysRevB.81.052405.
40. Lohmann A., Schmidt H.-J., Richter J., Phys. Rev. B, 2014, 89, 052405, doi:10.1103/PhysRevB.89.052405.
41. Juhász Junger I., Ihle D., Richter J., Phys. Rev. B, 2009, 80, 052405, doi:10.1103/PhysRevB.80.052405.
42. Le Guillou J.C., Zinn-Justin J., Phys. Rev. B, 1980, 21, 3976, doi:10.1103/PhysRevB.21.3976.
43. Holm C., Janke W., Phys. Rev. B, 1993, 48, 936, doi:10.1103/PhysRevB.48.936.
44. Holm C., Janke W., Phys. Rev. Lett., 1997, 78, 2265, doi:10.1103/PhysRevLett.78.2265.
45. De Souza A.J.F., Costa U.M.S., Lyra M.L., Phys. Rev. B, 2000, 62, 8909, doi:10.1103/PhysRevB.62.8909.
46. Schmidt H.-J., Lohmann A., Richter J., Phys. Rev. B, 2011, 84, 104443, doi:10.1103/PhysRevB.84.104443.
47. Müller P., Lohmann A., Richter J., Derzhko O., (unpublished).

Спін-1/2 феромагнітник Гайзенберга на гратці пірохлору. Дослідження методом функцій Гріна

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Ми розглядаємо квантовий феромагнетик Гайзенберга на гратці пірохлору і обговорюємо властивості цієї спінової моделі при довільних температурах. Для цього ми використовуємо техніку функцій Гріна у наближених хаотичних фаз (наближених Тяблкова), а також лінійну теорію спінових хвиль і симуляції методом квантового Монте Карло. Для підтвердження наших результатів ми порівнюємо їх з отриманими нами ефективними способами. Нарешті, ми співставляємо наші результати з відповідними для простої кубічної гратки, адже обидві гратки ідентичні у середньопольовій картині. Ми показуємо, що теплові флукутації ефективніші у випадку гратки пірохлору (ефект фрустрації при скінченних температурах). На основі результатів можна використовувати для згуртування експериментальних даних для матеріалів, у яких реалізуються феромагнетики на гратці пірохлору.

Ключові слова: квантові спінові системи, модель Гайзенберга, гратка пірохлору, функції Гріна, наближення хаотичних фаз
